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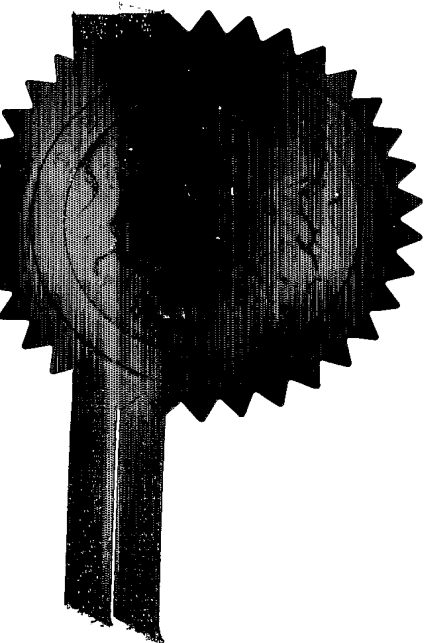
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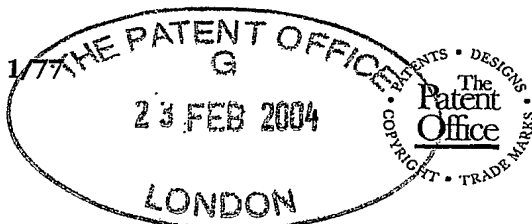


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Andrew Gersey

Dated 31 March 2005





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2. Patent application number 0403992.1
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Patents ADP number (if you know it)
If the applicant is a corporate body, give the country/state of its incorporation

ISIS INNOVATION LIMITED
Ewert House
Ewert Place
Summertown
Oxford
OX2 7SG
United Kingdom

08618878002

4. Title of the invention
OXIDATION BY HYDROGEN PEROXIDE

5. Name of your agent (if you have one)
"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

J. A. KEMP & CO.
14 South Square
Gray's Inn
London
WC1R 5JJ

Patents ADP number (if you know it) 26001

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Continuation sheets of this form	-
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11. I/We request the grant of a patent on the basis of this application.

Signature(s)

J. A. Kemp & Co

J.A. KEMP & CO.

Date 23 February 2004

12. Name, daytime telephone number and e-mail address, if any, of person to contact in the United Kingdom

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OXIDATION BY HYDROGEN PEROXIDE

Field of the Invention

The invention relates to a method of carrying out an oxidation reaction.

Background of the Invention

Monooxygenase enzymes catalyse the oxidation of a very wide range of substrates. In order to catalyse the reaction, a monooxygenase enzyme generally requires a cofactor and at least one electron-transfer partner protein (reductase).

However, monooxygenase enzymes are capable of using hydrogen peroxide (H_2O_2) as an oxidizing agent because it acts as a source of dioxygen and two electrons. The use of H_2O_2 to drive oxidation reactions is known as the "peroxide shunt".

Summary of the invention

Monooxygenase enzymes generally have a high K_m for H_2O_2 , (such as about 20mM) in comparison to peroxidase enzymes. As a result, high concentrations of H_2O_2 are required for appreciable levels of activity of a monooxygenase enzyme when the oxidation reaction is performed using the peroxide shunt. For example, the initial rate of monooxygenase activity using 50mM H_2O_2 is far below that when the natural co-factor, NAD(P)H, is used as with the physiological electron-transfer partners.

The invention provides a more efficient method of carrying out an oxidation reaction using the peroxide shunt by reducing the oxidative damage that occurs to the monooxygenase enzyme by not allowing excess levels of H_2O_2 to be present whilst the reaction is carried out.

Simultaneous production of H_2O_2 at a rate less than or equal to the rate at which it is used in an oxidation reaction catalysed by monooxygenase results in improved efficiency of the oxidation reaction and an increase in the product yield. Various methods may be used to produce H_2O_2 at the required rate, such as use of an electrochemical reaction, an enzyme or a precursor.

Accordingly, the present invention provides a method of carrying out an oxidation reaction catalysed by a monooxygenase enzyme and using hydrogen peroxide as an oxidant, in which reaction a low level of oxidation damage of the monooxygenase occurs, said method comprising producing the hydrogen peroxide

simultaneously with the oxidation reaction, wherein the hydrogen peroxide is produced at a rate less than or equal to the rate at which it is used in the reaction.

The present invention also provides a method of carrying out an oxidation reaction catalysed by a monooxygenase enzyme and using hydrogen peroxide as an oxidant, in which reaction a low level of oxidation damage of the monooxygenase occurs, said method comprising carrying out the reaction in the presence of an H₂O₂ or hydroxyl radical sequestering agent that controls the H₂O₂ or hydroxyl radical concentration.

Description of the Sequences

SEQ ID NO: 1 shows the nucleotide sequence of cytochrome P450Cam from *Pseudomonas putida*.

SEQ ID NO: 2 shows the amino acid sequence of cytochrome P450Cam from *Pseudomonas putida*.

SEQ ID NO: 3 shows the nucleotide sequence of cytochrome P450BM-3 from *Bacillus megaterium*.

SEQ ID NO: 4 shows the amino acid sequence of cytochrome P450 BM-3 from *Bacillus megaterium*. The first 472 amino acid residues form the heme domain. The last 585 amino acid residues form the reductase domain. All 1048 amino acid residues form the holoenzyme.

The convention in the art, which is adopted herein, is to refer to a mutant with reference to the native amino acid residue at a position in the sequence, followed by the amino acid at that position in the mutant, e. g., F87 refers to the phenylalanine at position 87 in the wild-type sequence, and F87A refers to the phenylalanine at position 87 in the wild-type sequence which has been changed to alanine in the variant. The numbering of the amino acid residues starts with the amino acid residue following the initial methionine residue.

Mutants used in Examples were F87A (single mutation; SEQ ID NOs: 5 and 6) and F87V L188Q A74G (triple mutation; SEQ ID NOs: 7 and 8).

SEQ ID NO: 5 shows the amino acid sequence of the F87A mutant of cytochrome P450BM-3 from *Bacillus megaterium*.

SEQ ID NO: 6 shows the nucleotide sequence of of the F87A mutant of cytochrome P450BM-3 from *Bacillus megaterium*.

SEQ ID NO: 7 shows the amino acid sequence of the F87V L188Q A74G

mutant of cytochrome P450BM-3 from *Bacillus megaterium*.

SEQ ID NO: 8 shows the nucleotide sequence of of the F87V L188Q A74G mutant of cytochrome P450BM-3 from *Bacillus megaterium*.

5 SEQ ID NO: 9 shows the nucleotide sequence of subunit 1 of B-276 alkene epoxidase from *Nocardia coralline*.

SEQ ID NO: 10 shows the amino acid sequence of subunit 1 of B-276 alkene epoxidase from *Nocardia coralline*.

SEQ ID NO: 11 shows the nucleotide sequence of subunit 2 of B-276 alkene epoxidase from *Nocardia coralline*.

10 SEQ ID NO: 12 shows the amino acid sequence of subunit 2 of B-276 alkene epoxidase from *Nocardia coralline*.

SEQ ID NO: 13 shows the nucleotide sequence of the alpha subunit of Py2 alkene monooxygenase from *Xanthobacta* sp.

15 SEQ ID NO: 14 shows the amino acid sequence of the alpha subunit of Py2 alkene monooxygenase from *Xanthobacta* sp.

SEQ ID NO: 15 shows the nucleotide sequence of the beta subunit of Py2 alkene monooxygenase from *Xanthobacta* sp.

SEQ ID NO: 16 shows the amino acid sequence of the beta subunit of Py2 alkene monooxygenase from *Xanthobacta* sp.

20 SEQ ID NO: 17 shows the nucleotide sequence of the gamma subunit of Py2 alkene monooxygenase from *Xanthobacta* sp.

SEQ ID NO: 18 shows the amino acid sequence of the gamma subunit of Py2 alkene monooxygenase from *Xanthobacta* sp.

25 SEQ ID NO: 19 shows the nucleotide sequence of the alpha subunit of soluble methane monooxygenase from *Methylococcus capsulatas*.

SEQ ID NO: 20 shows the amino acid sequence of the alpha subunit of soluble methane monooxygenase from *Methylococcus capsulatas*.

SEQ ID NO: 21 shows the nucleotide sequence of the beta subunit of soluble methane monooxygenase from *Methylococcus capsulatas*.

30 SEQ ID NO: 22 shows the amino acid sequence of the beta subunit of soluble methane monooxygenase from *Methylococcus capsulatas*.

SEQ ID NO: 23 shows the nucleotide sequence of the gamma subunit of soluble methane monooxygenase from *Methylococcus capsulatas*.

SEQ ID NO: 24 shows the amino acid sequence of the gamma subunit of

soluble methane monooxygenase from *Methylococcus capsulatus*.

SEQ ID NO: 25 shows the nucleotide sequence of GPO1 alkane hydroxylase (*AlkB* gene) from *Pseudomonas oleovorans*.

5 SEQ ID NO: 26 shows the amino acid sequence of GPO1 alkane hydroxylase from *Pseudomonas oleovorans*.

SEQ ID NO: 27 shows the nucleotide sequence of the alpha subunit of toluene 2-monooxygenase from *Burkholderia cepacia*.

SEQ ID NO: 28 shows the amino acid sequence of the alpha subunit of toluene 2-monooxygenase from *Burkholderia cepacia*.

10 SEQ ID NO: 29 shows the nucleotide sequence of the beta subunit of toluene 2-monooxygenase from *Burkholderia cepacia*.

SEQ ID NO: 30 shows the amino acid sequence of the beta subunit of toluene 2-monooxygenase from *Burkholderia cepacia*.

15 SEQ ID NO: 31 shows the nucleotide sequence of the gamma subunit of toluene 2-monooxygenase from *Burkholderia cepacia*.

SEQ ID NO: 32 shows the amino acid sequence of the gamma subunit of toluene 2-monooxygenase from *Burkholderia cepacia*.

SEQ ID NO: 33 shows the nucleotide sequence of phenol hydroxylase (*pheA*) gene from *Bacillus stearothermophilus*.

20 SEQ ID NO: 34 shows the amino acid sequence of phenol hydroxylase gene from *Bacillus stearothermophilus*.

SEQ ID NO: 35 shows the nucleotide sequence of stearoyl-ACP desaturase from *Helianthus annuus*.

25 SEQ ID NO: 36 shows the amino acid sequence of stearoyl-ACP desaturase from *Helianthus annuus*.

Detailed description of the Invention

30 It is to be understood that this invention is not limited to particular embodiments. It is also to be understood that different applications of the disclosed methods may be tailored to the specific needs in the art. It is also to be understood that the terminology used herein is for the purpose of describing particular embodiments of the invention only, and is not intended to be limiting.

In addition as used in this specification and the appended claims, the singular forms "a", "an", and "the" include plural referents unless the content clearly

dictates otherwise. Thus, for example, reference to "a substrate" includes two or more substrates, reference to "an enzyme" includes reference to two or more enzymes, and the like.

5 All publications, patents and patent applications cited herein, whether supra or infra, are hereby incorporated by reference in their entirety.

The methods of the invention enable the oxidation of a variety of substrates. Such substrates include, but are not limited to, alkanes, aromatic compounds, terpenoid compounds, alkenes and fatty acids.

10 Suitable alkanes include, but are not limited to, methane, ethane, propane, butane, pentane, hexane, heptane, *n*-octane, *n*-nonane, *n*-decane, *n*-dodecane and *n*-hexadecane. The oxidation of alkanes produces alcohols. The oxidation of methane to methanol is technologically and economically very important. The medium-chain alcohols (e.g. *n*-octanol) are synthetic intermediates while the longer chain alcohols (e.g. *n*-dodecanol) are used for the synthesis of fatty acid derivatives.

15 Suitable aromatic compounds include, but are not limited to, benzene, toluene, xylene, chlorobenzene, phenol and substituents thereof. The phenolic and catecholic products are used in the synthesis of fragrance and flavour compounds.

20 Suitable terpenoid compounds include, but are not limited to, monoterpenes such as limonene, pinene, terpinene, and ocimene, sesquiterpenes such as valencene and aromadendrene and triterpenes which include the steroidal compounds. The products are intermediates for synthesis, fine fragrance and flavouring chemicals and pharmaceuticals.

25 Suitable alkenes include, but are not limited to, simple molecules such as propene, hex-1-ene, hex-2-ene, and styrene, and carbon-carbon double bonds in complex molecules. Selective epoxidation of alkenes to a single enantiomer is very important in synthesis. Optically pure propene oxide and styrene oxide are very useful intermediates in synthesis.

Hydroxylated fatty acids are precursors to polymers.

30 *Monooxygenase enzyme*

The enzyme used to carry out an oxidation reaction according to the invention is a monooxygenase enzyme. A person skilled in the art can determine whether an enzyme is a monooxygenase enzyme using standard techniques in the art. Typically, the prosthetic groups may be characterised using protein crystallography,

especially for non-heme iron enzymes because they generally do not have chromophores. Otherwise, a person skilled in the art will typically use sequence alignment, looking for conserved motifs such as the active site, and iron content as well as subunit composition.

5 The monooxygenase enzyme preferably has a K_m for H_2O_2 of at least 15nM, at least 20nM, at least 25nM, at least 30nM, at least 35nM, at least 40nM, at least 45nM or at least 50nM.

 Examples of monooxygenase enzymes include, but are not limited to, cytochrome P450 monooxygenases and non-heme di-iron monooxygenase enzymes.

10 Suitable non-heme di-iron monooxygenase enzymes include, but are not limited to methane monooxygenase (Colby *et al.*, Biochem. J., 1977; 165: 395-402; Dalton, Adv. Appl. Microbiol., 1980; 26: 71-87; Fox *et al.*, J. Biol. Chem., 1989; 264: 10023-10033; Fox *et al.*, Methods Enzymol., 1990; 188: 191-202; McDonald *et al.*, Appl. Environ. Microbiol., 1997; 63: 1898-1904), alkane hydroxylase (van Beilen *et al.*, Enzyme Microb. Technol., 1994; 16: 904-911), toluene monooxygenase (Luykx *et al.*, Biochem. Biophys. Res. Commun., 2003; 312: 373-379; Pikus *et al.*, Biochemistry, 1996; 35: 9106-9119; Newman & Wackett, Biochemistry, 1995; 34: 14066-14076), alkene monooxygenase (Gallagher *et al.*, Eur. J. Biochem., 1997; 247: 635-641; Lange & Que, Curr. Opin. Chem. Biol., 1998; 2: 159-172; Zhou *et al.*, FEBS Lett., 1998; 430: 181-185), phenol monooxygenase (Divari *et al.*, Eur. J. Biochem., 2003; 270: 2244-2253) and steroid desaturase (Shanklin *et al.*, Biochemistry, 1994; 33: 12787-12794). The non-heme di-iron monooxygenase enzymes are typically of eukaryotic or prokaryotic origin and preferably of bacterial, fungal, yeast, plant or animal origin. Preferred sequences are shown in SEQ ID NOs:

20 1 to 36.

 The enzyme used in the methods of the invention is preferably a cytochrome P450 enzyme, typically of eukaryotic or prokaryotic origin. Cytochrome P450 monooxygenases are typically characterised by a 446-450 nm heme Soret band for the ferrous-carbon monoxide complex. The enzyme is generally of bacterial, fungal,

30 yeast, plant or animal origin, and thus may be from a bacterium of the genus *Pseudomonas*. The enzyme may be a naturally-occurring form of P450, such as P450_{cam}, P450_{BM-3} from *Bacillus megaterium*, P450_{terp} from *Pseudomonas sp.*, P450_{eryF} from *Saccharopolyspora erythraea* and also P450 105 D1 (CYP105) from *Streptomyces griseus* strains.

Alternatively, the enzyme may be a mutant of a naturally-occurring form of P450. The mutants retain the essential biological activity of the naturally-occurring enzyme, namely the ability to catalyse an oxidation reaction using H_2O_2 . The mutant may have one or more mutations in the active site of the enzyme.

5 An amino acid 'in the active site' is one which lines or defines the site in which the substrate is bound during catalysis or one which lines or defines a site through which the substrate must pass before reaching the catalytic site. Therefore such an amino acid typically interacts with the substrate during entry to the catalytic site or during catalysis. Such an interaction typically occurs through an electrostatic
10 interaction (between charged or polar groups), hydrophobic interaction, hydrogen bonding or van der Waals forces.

The amino acids in the active site can be identified by routine methods to those skilled in the art. These methods include labelling studies in which the enzyme is allowed to bind a substrate which modifies ('labels') amino acids which contact
15 the substrate. Alternatively the crystal structure of the enzyme with bound substrate can be obtained in order to deduce the amino acids in the active site.

The monooxygenase enzyme may have 1, 2, 3, 4, 5 to 10, 10 to 20 or more other mutations, such as substitutions, insertions or deletions. Amino acid
20 substitutions may be made to the amino acid sequence of a naturally-occurring enzyme, for example from 1, 2, 3, 4 or 5 to 10, 20 or 30 substitutions. Conservative substitutions may be made, for example, according to Table 1. Amino acids in the same block in the second column and preferably in the same line in the third column may be substituted for each other:

25 **Table 1 – Conservative amino acid substitutions**

NON-AROMATIC	Non-polar	G A P
		I L V
	Polar – uncharged	C S T M
		N Q
	Polar - charged	D E
		H K R
AROMATIC		H F W Y

The mutations may be in the active site or outside the active site. Typically the mutations are in the 'second sphere' residues which affect or contact the position or orientation of one or more of the amino acids in the active site. The insertion is typically at the N and/or C terminal and thus the enzyme may be part of a chimeric protein. The deletion typically comprises the deletion of amino acids which are not involved in catalysis, such as those outside the active site (thus the enzyme is a mutated fragment of a naturally occurring enzyme). The monooxygenase enzyme may thus comprise only those amino acids which are required for oxidation activity.

The mutation in the active site typically alters the position and/or conformation of the substrate when it is bound in the active site. The mutation may make the site on the substrate which is to be oxidized more accessible to the heme group. Thus the mutation may be a substitution to an amino acid which has a smaller or larger, or more or less polar, side chain.

The mutations typically increase the stability of the protein, or make it easier to purify the protein. They typically prevent the dimerisation of the protein, typically by removing cysteine residues from the protein (e.g. by substitution of cysteine at position 334 of P450_{cam}, or at an equivalent position in a homologue, preferably to alanine). They typically allow the protein to be prepared in soluble form, for example by the introduction of deletions or a poly-histidine tag, or by mutation of the N-terminal membrane anchoring sequence. The mutations typically inhibit protein oligomerisation, such as oligomerisation arising from contacts between hydrophobic patches on protein surfaces.

The mutations may affect the manner in which the enzyme utilises H₂O₂ and thereby improve the efficiency of the reaction. For example, mutants of the P450 enzyme from *Pseudomonas putida* hydroxylate naphthalene through the "peroxide shunt" with more than a 20-fold increase in the activity of the enzyme (Joo *et al.*, Nature, 1999; 399(6737): 636-637). In addition, replacement of all the methionine residues of the heme domain of P450_{BM-3} with norleucine results in a two-fold increase in the peroxxygenase activity of the enzyme (Cirino *et al.*, Biotechnol. Bioeng., 2003; 83(6): 729-734). Furthermore, direct evolution studies to find mutants of enzymes more resistant to peroxide (Cirino & Arnold, Angew. Chem. Int. Ed., 2003; 42: 3299-3301).

Thus the mutant enzyme is typically at least 70% homologous to a naturally occurring enzyme on the basis of amino acid identity.

A mutant protein (i.e. described as being a mutant of another protein) mentioned herein is typically at least 70% homologous to the relevant protein or at least 80 or 90% and more preferably at least 95%, 97% or 99% homologous thereto over at least 20, preferably at least 30, for instance at least 40, 60 or 100 or more contiguous amino acids. The contiguous amino acids may include the active site. This homology may alternatively be measured not over contiguous amino acids but over only the amino acids in the active site.

Homology can be measured using known methods. For example the UWGCG Package provides the BESTFIT program which can be used to calculate homology (for example used on its default settings) (Devereux *et al* (1984) *Nucleic Acids Research* **12**, p387-395). The PILEUP and BLAST algorithms can be used to calculate homology or line up sequences (typically on their default settings), for example as described in Altschul S. F. (1993) *J Mol Evol* 36:290-300; Altschul, S. F. *et al* (1990) *J Mol Biol* 215:403-10.

Software for performing BLAST analyses is publicly available through the National Center for Biotechnology Information (<http://www.ncbi.nlm.nih.gov/>). This algorithm involves first identifying high scoring sequence pair (HSPs) by identifying short words of length W in the query sequence that either match or satisfy some positive-valued threshold score T when aligned with a word of the same length in a database sequence. T is referred to as the neighbourhood word score threshold (Altschul *et al*, supra). These initial neighbourhood word hits act as seeds for initiating searches to find HSPs containing them. The word hits are extended in both directions along each sequence for as far as the cumulative alignment score can be increased. Extensions for the word hits in each direction are halted when: the cumulative alignment score falls off by the quantity X from its maximum achieved value; the cumulative score goes to zero or below, due to the accumulation of one or more negative-scoring residue alignments; or the end of either sequence is reached. The BLAST algorithm parameters W, T and X determine the sensitivity and speed of the alignment. The BLAST program uses as defaults a word length (W) of 11, the BLOSUM62 scoring matrix (see Henikoff and Henikoff (1992) *Proc. Natl. Acad. Sci. USA* 89: 10915-10919) alignments (B) of 50, expectation (E) of 10, M=5, N=4, and a comparison of both strands.

The BLAST algorithm performs a statistical analysis of the similarity between two sequences; see e.g., Karlin and Altschul (1993) *Proc. Natl. Acad. Sci.*

USA 90: 5873-5787. One measure of similarity provided by the BLAST algorithm is the smallest sum probability ($P(N)$), which provides an indication of the probability by which a match between two nucleotide or amino acid sequences would occur by chance. For example, a sequence is considered similar to another
5 sequence if the smallest sum probability in comparison of the first sequence to the second sequence is less than about 1, preferably less than about 0.1, more preferably less than about 0.01, and most preferably less than about 0.001.

Mutants include fragments of the above-mentioned sequences. Such fragments retain monooxygenase activity. Fragments may be at least 300, at least
10 400 or at least 450 amino acids in length. Such fragments may be used to produce chimeric enzymes as described in more detail below.

Mutants also include chimeric proteins comprising fragments or portions of a naturally-occurring enzyme. One or more amino acids may be alternatively or additionally added to the polypeptides described above. An extension may be
15 provided at the N-terminus or C-terminus of the naturally-occurring enzyme or variant or fragment thereof. The extension may be quite short, for example from 1 to 10 amino acids in length. Alternatively, the extension may be longer. A carrier protein may be fused to an amino acid sequence described above. A fusion protein incorporating one of the enzymes described above can thus be used in the invention.

20 The naturally-occurring enzyme or mutant thereof may also be chemically-modified. A number of side chain modifications are known in the art and may be made to the side chains of the enzymes discussed above. Such modifications include, for example, glycosylation, phosphorylation, modifications of amino acids by reductive alkylation by reaction with an aldehyde followed by reduction with
25 NaBH_4 , amidination with methylacetimidate or acylation with acetic anhydride. The modification is preferably glycosylation.

The mutations discussed herein are generally introduced into the enzyme by using methods known in the art, such as site directed mutagenesis of the enzyme, PCR and gene shuffling methods or by the use of multiple mutagenic
30 oligonucleotides in cycles of site-directed mutagenesis. Thus the mutations may be introduced in a directed or random manner. The mutagenesis method thus produces one or more polynucleotides encoding one or more different mutants. Typically a library of mutant oligonucleotides is produced which can be used to produce a library of mutant enzymes.

The enzyme may be made synthetically or by recombinant means using methods known in the art. The amino acid sequence of the monooxygenase enzyme may be modified to include non-naturally occurring amino acids or to increase the stability of the enzyme. When the enzyme is produced by synthetic means, such amino acids may be introduced during production. The proteins or peptides may also be modified following either synthetic or recombinant production.

The enzyme may also be produced using D-amino acids. In such cases the amino acids will be linked in reverse sequence in the C to N orientation. This is conventional in the art for producing such proteins or peptides.

The enzyme may be produced in a cell by *in situ* expression of the polypeptide from a recombinant expression vector. The expression vector optionally carries an inducible promoter to control the expression of the polypeptide. The enzyme may be produced in large scale following purification by any protein liquid chromatography system after recombinant expression. Preferred protein liquid chromatography systems include FPLC, AKTA systems, the Bio-Cad system, the Bio-Rad BioLogic system and the Gilson HPLC system.

Oxidation reaction

The methods of the invention concerns carrying out a high efficiency oxidation reaction catalysed by a monooxygenase enzyme. A high efficiency oxidation reaction is a reaction that occurs without an appreciable reduction in the enzyme turnover or product yield or inactivation of the monooxygenase enzyme. Preferably, the monooxygenase enzyme displays at least 70%, at least 80%, at least 90%, at least 95% or 100% of the activity shown at the beginning of the reaction after 1 hour, 2 hours, 6 hours, 12 hours, 1 day, 2 days or 5 days.

Typically the methods of the invention are carried out *in vitro*, such as in a cell free system.

The reaction is driven by the "peroxide shunt". The reaction of the invention is carried out in the presence of the monooxygenase enzyme (a), the substrate (b) and H_2O_2 (c). The reaction is typically performed in aerobic conditions and does not require any cofactors. The production of (c) is discussed in more detail below. In this system the flow of electrons is typically: (c) \rightarrow (a) \rightarrow (b).

In the methods the concentration of (a) and (b) is typically from 10^{-8} to 10^{-2} M, preferably from 10^{-6} to 10^{-4} M. Typically the ratio of concentrations of (a): (b) is

from 0.1:10 to 1:10, preferably from 1:0.5 to 1:2, or from 1:0.8 to 1:1.2. Preferably, the concentration of (b) is greater than the concentration of (a). The preferred concentration of (a) is that which when reacted with substrate will generate sufficient product to be detected by available analytical methods e.g. GC, HPLC. This is
5 typically of the order of μM quantities.

Generally the methods are carried out at a temperature and/or pH at which the monooxygenase enzyme is functional, such as when the enzyme has at least 20%, 50%, 80% or more of peak activity. Typically the pH is from 2 to 11, such as from 5 to 9 or from 6 to 8, preferably from 7 to 7.8 or 7.4. The pH can be maintained using
10 a suitable buffering agent such as phosphate or acetate based systems. Typically the temperature is from 0 to 80°C, such as from 25 to 75°C, from 30 to 60°C or from 50°C to 80°C. Preferably, the temperature is from 20 to 40°C.

Typically in the methods at least 20 turnovers/min occur, such as at least 50, 100, 200, 300, 500 or more turnovers (turnover is measured as nanomoles of product
15 formed per nanomole of enzyme).

Typically, the rate of H_2O_2 production is less than or equal to 1, 2 or 3 μg per min per mg of monooxygenase enzyme. Typically, the concentration of H_2O_2 throughout the reaction is less than or equal to 0.1, 0.5 or 1mM. Typically, the reaction continues for at least 60 minutes, at least 240 minutes, at least 6 hours or at
20 least 10 hours.

The methods of the invention may be carried out in the monooxygenase substrate if it is a liquid under the reaction conditions. The methods of the invention may also be conducted in a solvent. Suitable solvents include, but are not limited to, water, aqueous buffer solutions mixed water/organic and aqueous buffer/organic
25 solvent systems. Preferably, the organic solvent is a hydrocarbon such as hexane, benzene, acetonitrile, lower aliphatic alcohols, ketones and dioxane, dimethylformamide and dimethylsulphoxide and mixtures thereof. The solvent is typically one in which the reagents and products are highly soluble and one that maintains the stability and activity of the monooxygenase enzyme.

30 The reaction may be carried out in a homogenous system with all the components in solution. Typically, the monooxygenase enzyme and substrate are mixed together in a suitable solvent in a stirred tank reactor and the reaction is conducted in batch, semi-batch or continuous mode.

Alternatively, the monooxygenase enzyme may be immobilized on a suitable solid support, such as silica, prior to carrying out the method of the invention. An immobilized monooxygenase enzyme can be packed into a fixed bed reactor and the substrate passed over the enzyme. In one embodiment, the enzyme
5 producing the H_2O_2 (discussed in more detail below) may be immobilized on the same or different material as the monooxygenase enzyme. Procedures for immobilizing enzymes are known in the art. Examples of such procedures include, but are not limited to, covalent coupling to insoluble organic or inorganic supports, entrapment in gels and adsorption to ion exchange resins or other adsorbent
10 materials. (G. F. Bickerstaff ed., "Immobilization of Enzymes and Cells," Humana Press, Totowa, New Jersey, 1997).

In a further embodiment, a membrane on the "entry" side admits the substrate slowly from the "reactant" side and then a hydrophilic membrane on the "exit" side allows hydrophilic compounds to flow out to the "product" side of the
15 flow reaction cell. In this case the H_2O_2 may be generated outside the membrane and allowed to flow through the membrane to the mobile or immobile enzyme.

In one embodiment, H_2O_2 is preferably produced by one of the methods discussed in more detail below. In another embodiment, a H_2O_2 or hydroxyl radical sequestering agent is used to sequester excess H_2O_2 or hydroxyl radical during the
20 oxidation reaction. The sequestering agent may be a chelating agent. In one embodiment, the chelating agent is EDTA. The EDTA inhibits production of the hydroxyl radical, for example, produced by the reaction of trace amounts of iron (or copper) with the H_2O_2 .

25 *H_2O_2 production by an electrochemical reaction*

The H_2O_2 may be produced in the method of the invention by an electrochemical reaction. An electrochemical reaction is generally a means for introducing a current to a liquid, preferably a solution. An electrochemical reaction is typically an oxidation or reduction reaction that takes place at an electrode through
30 which a current flows. An electrode is a solid capable of conducting electricity, typically carbon-based or metallic, leading to an external source or sink which is in contact with the liquid, preferably a solution. The electrode may be either positively charged (cathode) or negatively charged (anode). Two or more electrodes may form an electrochemical cell from which an external wire can lead from each electrode to

an external electrical device. An oxidation or reduction reaction takes place at one electrode, while a redox reaction can take place either in an electrochemical cell or directly in the liquid.

- Production of H_2O_2 using an electrochemical reaction is energy efficient.
- 5 H_2O_2 is typically produced by the controlled electrochemical reduction of molecular oxygen to hydrogen peroxide. The surface area and the overpotential of the cathode are key considerations for the two-electron reduction of molecular oxygen to hydrogen peroxide. Typically, carbon-based cathodes are used and they may be modified with a compound known to lower the overpotential for this reaction.
- 10 Electrode materials and modifiers which will perform this task effectively and efficiently are well known in the art. The reduction of O_2 , and hence production of hydrogen peroxide, can typically be controlled by the potential applied to the cathode. The potential applied to the cathode will vary depending on the cathode and any modifications to the cathode made.
- 15 The electrochemical reaction used in the method of the invention may be the sonoelectrochemical reduction of dioxygen. This method is well known in the art (Compton *et al.*, *Electroanalysis*, 1997; 9(7): 509-522).

H_2O_2 production by an enzyme

- 20 The H_2O_2 may be produced in the method of the invention by an enzyme. The enzyme is preferably an oxidase. Examples of suitable oxidases include, but are not limited to, glucose oxidase (E.C. 1.1.3.4), secondary-alcohol oxidase (E.C. 1.1.3.18), methanol oxidase (E.C. 1.1.3.31), oxalate oxidase (E.C. 1.2.3.4), aryl-aldehyde oxidase (E.C. 1.2.3.9), carbon monoxide oxidase (E.C. 1.2.3.10), amine
- 25 oxidase (E.C. 1.4.3.4), ethanolamine oxidase (E.C. 1.4.3.8), nitroethane oxidase (E.C. 1.7.3.1) and sulfite oxidase (E.C. 1.8.3.1). Glucose oxidase (E.C. 1.1.3.4) catalyzes the conversion of D-glucose to D-glucono-1,5-lactone and H_2O_2 . Secondary-alcohol oxidase (E.C. 1.1.3.18) catalyzes the conversion of a secondary alcohol to a ketone and H_2O_2 . Methanol oxidase (E.C. 1.1.3.31) catalyzes the
- 30 conversion of methanol to formaldehyde and H_2O_2 . Oxalate oxidase (E.C. 1.2.3.4) catalyzes the conversion of oxalate to carbon dioxide and H_2O_2 . Aryl-aldehyde oxidase (E.C. 1.2.3.9) catalyzes the conversion of an aromatic aldehyde to an aromatic acid and H_2O_2 . Carbon monoxide oxidase (E.C. 1.2.3.10) catalyzes the conversion of CO and H_2O to carbon dioxide and H_2O_2 . Amine oxidase (E.C.

1.4.3.4) catalyzes the conversion of RCH_2NH_2 and H_2O to RCHO and NH_3 and H_2O_2 . Ethanolamine oxidase (E.C. 1.4.3.8) catalyzes the conversion of ethanolamine and H_2O to glycolaldehyde and H_2O_2 . Nitroethane oxidase (E.C. 1.7.3.1) catalyzes the conversion of nitroethane and H_2O to acetaldehyde and H_2O_2 . Sulfite oxidase (E.C. 1.8.3.1) catalyzes the conversion of sulfite and H_2O_2 to sulfate and H_2O_2 . The oxidase may be purchased commercially (e.g., glucose oxidase). Alternatively, the oxidase can be extracted from known microorganisms using procedures known in the art.

The substrate for the oxidase will be well known in the art. In addition to the substrate, the reaction to produce H_2O_2 will also involve water. Typically, a H_2O_2 -activating metal is also included in the reaction. Suitable metals include, but are not limited to, cerium, chromium, cobalt, copper, iron, manganese, molybdenum, silver, titanium, tungsten, vanadium and mixtures thereof. Metallosilicates containing the above metals can be prepared and used in the method of the invention. The procedure for producing such metallosilicates is known in the art (Neumann *et al.*, Journal of Catalysis, 1997; 166: 206-127). The metallosilicate is preferably tetrahedrally coordinated titanium such as silicalite-1 (TS-1), silicalite-2 (TS-2), zeolite-beta, silicon analogs of ZSM-48 and MCM-4 1. (Murugavel and Roesky, Angew. Chem. Int. Ed. Engl., 1997; 36(5): 477-479).

In a preferred embodiment of the invention, the metal-containing solid or metallosilicate is used as a support upon which the H_2O_2 -producing enzyme is immobilized. In another preferred embodiment, the monooxygenase enzyme is also immobilized on the same or different metallosilicate support.

Preferably, the oxidase is first mixed with the other reaction components and then the reaction is initiated by addition of the oxidase substrate. For example, the monooxygenase enzyme, monooxygenase enzyme substrate and oxidase are all mixed and then the oxidase enzyme is added. In a preferred embodiment, P450_{BM3} , octane and glucose oxidase are mixed together and then glucose added. Control of H_2O_2 generation can typically be accomplished by controlling the rate at which the oxidase substrate is added.

H_2O_2 production by a precursor

The H_2O_2 may be produced in the method of the invention by a precursor. The generation of H_2O_2 by the addition of a precursor to water is well known in the

art. Precursors include, but are not limited to, salts of perborate, salts of percarbonate, salts of perphosphate and peroxyxynitrite. Preferred precursors are sodium salts. The H_2O_2 -producing properties of the precursor may be enhanced by using a compound such as tetraacetythylenediamine. The amount of precursor added to the solution containing the monooxygenase enzyme and substrate is such to maximise the enzymatic reaction with the substrate and to minimise the deactivation of the enzyme by H_2O_2 . Preferably the concentration of H_2O_2 produced does not exceed the K_m value for the enzyme but is sufficient to generate the enzyme reactive species.

10

Examples

Example 1

In this experiment, octane was reacted with electrochemically generated H_2O_2 in the presence of P450_{BM3} heme domain. The experiment was performed at room temperature with a three-electrode configuration in a 100 mL glass beaker. The reticulated vitreous carbon (RVC) cathode, platinum gauze anode and Ag/AgCl reference electrode were contained in the one vessel. The RVC cathode was briefly immersed in a 1 mM 2-aminoanthraquinone ethanolic solution before being removed and allowed to dry in air. The reaction solution contained aqueous Tris buffer (50 mM, pH 7.4) saturated with oxygen, 0.2 M KCl, 0.5 mM octane, and 3 μM P450_{BM3} F87V L188Q A74G heme domain. The reaction solution was stirred to equilibrate (5-10 minutes) and then a potential of -0.55 V vs Ag/AgCl was applied for 2 hours and the solution stirred continuously throughout. GC analysis revealed the presence of the solvent chloroform, octane, 2-, 3- and 4-octanol and the internal standard 1-nonanol. The relative proportion of 2, 3 & 4-octanol was 1:1.1:0.7. The total concentration of octanols formed was 141 μM , representing a turnover per enzyme of 47.

A similar experiment was performed with 1.43 μM wild-type P450_{BM3} heme domain. The total concentration of octanols formed was 8.4 μM , representing a turnover per enzyme of 6. The relative proportion of 2, 3 & 4-octanol in this case was 1:1.7:2.0.

30

Example 2

In this experiment, octane was reacted with enzymatically generated H_2O_2 in the presence of P450_{BM3} holoenzyme. Into a glass vial was added a solution (total volume 5 mL) consisting of aqueous Tris buffer (50 mM, pH 7.4), 0.5 mM octane, 1.6 μM P450_{BM3} F87V L188Q A74G holoenzyme and glucose oxidase (1.5 U). After equilibration (5 mins), the reaction was initiated by addition of glucose (1×10^{-6} moles). Successive additions of glucose (1×10^{-6} moles) were made every 5 minutes up to 1 hour (total of 12 additions equivalent to 1.2×10^{-5} moles). The reaction was stirred continuously during this time and stopped after 1.5 hours. GC analysis revealed the presence of the solvent chloroform, octane, 2-, 3- and 4-octanol and the internal standard 1-nonanol. The relative proportion of 2, 3 & 4-octanol was 1:1.1:0.8. The total concentration of octanols formed was 17 μM , representing a turnover per enzyme of 10.

Example 3

In this experiment, octane was reacted with H_2O_2 derived from sodium perborate, in the presence of P450_{BM3} holoenzyme. Into a glass vial was added a solution (total volume 5 mL) consisting of aqueous Tris buffer (40 mM, pH 7.4), 0.5 mM octane, and 1.3 μM P450_{BM3} F87V L188Q A74G holoenzyme. After equilibration (5 mins), the reaction was initiated by addition of $\text{NaBO}_3 \cdot 4\text{H}_2\text{O}$ (1×10^{-4} moles) and stirred continuously for 1 hour. GC analysis revealed the presence of the solvent chloroform, octane, 2-, 3- and 4-octanol and the internal standard 1-nonanol. The relative proportion of 2, 3 & 4-octanol was 1:1.8:1.1. The total concentration of octanols formed was 77 μM , representing a turnover per enzyme of 59.

For Examples 1 to 3, no octanol products were observed when the P450 enzyme was absent from the solution.

Example 4

In this experiment, pinene was reacted with H_2O_2 derived from sodium perborate, in the presence of P450_{BM3} heme domain. Into a glass vial was added a solution (total volume 5 mL) consisting of aqueous Tris buffer (40 mM, pH 7.4), 0.63 mM pinene, and 3.7 μM wild-type P450_{BM3} heme domain. After equilibration (5 mins), the reaction was initiated by addition of 7.8×10^{-6} moles $\text{NaBO}_3 \cdot 4\text{H}_2\text{O}$ and

stirred continuously for 1 hour. GC analysis revealed the presence of *cis/trans* 2,3-epoxides (32%), (+)-*trans*-verbenol (16%), (+)-*cis*-verbenol (6%), (+)-verbenone/(+)-myrtenol (13%), myrtenal (4%), as well as unidentified further oxidation products (29%). The total concentration of products formed was 80 μ M, representing a turnover per enzyme of 22.

Example 5

In this experiment, phenol monooxygenase is reacted with phenol in the presence of with H_2O_2 generated by sodium perborate. Into a glass vial is added a solution (total volume 5 mL) consisting of aqueous Tris buffer (40 mM, pH 7.4), 0.63 mM phenol, and 3.7 μ M wild-type phenol monooxygenase. After equilibration (5 mins), the reaction is initiated by addition of 7.8×10^{-6} moles $NaBO_3 \cdot 4H_2O$ and stirred continuously for 1 hour. GC analysis reveals the presence of oxidation products.

Example 6

In this experiment, P450_{BM3} is reacted with palmitic acid in the presence of H_2O_2 generated by glucose oxidase. Into a glass vial is added a solution (total volume 5 mL) consisting of aqueous Tris buffer (50 mM, pH 7.4), 0.5 mM palmitic acid, 1.6 μ M P450_{BM3} holoenzyme and glucose oxidase (1.5 U). After equilibration (5 mins), the reaction is initiated by addition of glucose (1×10^{-6} moles). Successive additions of glucose (1×10^{-6} moles) are made every 5 minutes up to 1 hour (total of 12 additions equivalent to 1.2×10^{-5} moles). The reaction is stirred continuously during this time and stopped after 1.5 hours. GC analysis reveals the presence of oxidation products.

Example 7

Plant CYP74C is reacted with 13 S-hydroperoxylinolenic acid to form the compound 3Z-hexenal (a fragrance). The H_2O_2 is generated by sodium perborate. Into a glass vial is added a solution (total volume 5 mL) consisting of aqueous Tris buffer (40 mM, pH 7.4), 0.63 mM 13 S-hydroperoxylinolenic acid, and 3.7 μ M wild-type plant CYP74C. After equilibration (5 mins), the reaction is initiated by addition of 7.8×10^{-6} moles $NaBO_3 \cdot 4H_2O$ and stirred continuously for 1 hour. GC analysis

reveals the presence of oxidation products.

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Ala Met Met Val Asp Ile Ala Val Gln Leu Val Gln Lys Trp Glu Arg	
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Leu Asn Ala Asp Glu His Ile Glu Val Pro Glu Asp Met Thr Arg Leu	
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Thr Leu Asp Thr Ile Gly Leu Cys Gly Phe Asn Tyr Arg Phe Asn Ser	
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Phe Tyr Arg Asp Gln Pro His Pro Phe Ile Thr Ser Met Val Arg Ala	
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Ala Tyr Asp Glu Asn Lys Arg Gln Phe Gln Glu Asp Ile Lys Val Met	
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Asn Asp Leu Val Asp Lys Ile Ile Ala Asp Arg Lys Ala Ser Gly Glu	
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Gln Ser Asp Asp Leu Leu Thr His Met Leu Asn Gly Lys Asp Pro Glu	
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Thr Gly Glu Pro Leu Asp Asp Glu Asn Ile Arg Tyr Gln Ile Ile Thr	
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Phe Leu Ile Ala Gly His Glu Thr Thr Ser Gly Leu Leu Ser Phe Ala	
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Leu Tyr Phe Leu Val Lys Asn Pro His Val Leu Gln Lys Ala Ala Glu	
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Glu Ala Ala Arg Val Leu Val Asp Pro Val Pro Ser Tyr Lys Gln Val	
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Lys Gln Leu Lys Tyr Val Gly Met Val Leu Asn Glu Ala Leu Arg Leu	
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Trp Pro Thr Ala Pro Ala Phe Ser Leu Tyr Ala Lys Glu Asp Thr Val	
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Leu Gly Gly Glu Tyr Pro Leu Glu Lys Gly Asp Glu Leu Met Val Leu	
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Ile Pro Gln Leu His Arg Asp Lys Thr Ile Trp Gly Asp Asp Val Glu	
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Glu Phe Arg Pro Glu Arg Phe Glu Asn Pro Ser Ala Ile Pro Gln His	
375 380 385	
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Ala Phe Lys Pro Phe Gly Asn Gly Gln Arg Ala Cys Ile Gly Gln Gln	
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Phe Ala Leu His	Glu Ala Thr Leu Val Leu Gly Met Met Leu Lys His	
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Ile Pro Leu Gly Gly Ile Pro Ser Pro Ser Thr Glu Gln Ser Ala Lys		
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Lys Val Arg Lys Lys Ala Glu Asn Ala His Asn Thr Pro Leu Leu Val		
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Ser Val Phe Gly Cys Gly Asp Lys Asn Trp Ala Thr Thr Tyr Gln Lys		
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Ser Ala Arg Ser Thr Arg His Leu Glu Ile Glu Leu Pro Lys Glu Ala		
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Ser Tyr Gln Glu Gly Asp His Leu Gly Val Ile Pro Arg Asn Tyr Glu		
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Gln Ile Arg Leu Glu Ala Glu Glu Glu Lys Leu Ala His Leu Pro Leu	
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gct aaa aca gta tcc gta gaa gag ctt ctg caa tac gtg gag ctt caa	3811
Ala Lys Thr Val Ser Val Glu Glu Leu Leu Gln Tyr Val Glu Leu Gln	
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Cys Pro Pro His Lys Val Glu Leu Glu Ala Leu Leu Glu Lys Gln Ala	
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 50 55 60
 Glu Ser Arg Phe Asp Lys Asn Leu Ser Gln Ala Leu Lys Phe Val Arg
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 Asp Phe Ala Gly Asp Gly Leu Phe Thr Ser Trp Thr His Glu Lys Asn
 85 90 95
 Trp Lys Lys Ala His Asn Ile Leu Leu Pro Ser Phe Ser Gln Gln Ala
 100 105 110
 Met Lys Gly Tyr His Ala Met Met Val Asp Ile Ala Val Gln Leu Val
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 Gln Lys Trp Glu Arg Leu Asn Ala Asp Glu His Ile Glu Val Pro Glu
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 Tyr Arg Phe Asn Ser Phe Tyr Arg Asp Gln Pro His Pro Phe Ile Thr
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 Ser Met Val Arg Ala Leu Asp Glu Ala Met Asn Lys Leu Gln Arg Ala
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 Asn Pro Asp Asp Pro Ala Tyr Asp Glu Asn Lys Arg Gln Phe Gln Glu
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 Asp Ile Lys Val Met Asn Asp Leu Val Asp Lys Ile Ile Ala Asp Arg
 210 215 220
 Lys Ala Ser Gly Glu Gln Ser Asp Asp Leu Leu Thr His Met Leu Asn
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 Gly Lys Asp Pro Glu Thr Gly Glu Pro Leu Asp Asp Glu Asn Ile Arg
 245 250 255

Tyr Gln Ile Ile Thr Phe Leu Ile Ala Gly His Glu Thr Thr Ser Gly
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 Leu Leu Ser Phe Ala Leu Tyr Phe Leu Val Lys Asn Pro His Val Leu
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 Gln Lys Ala Ala Glu Glu Ala Ala Arg Val Leu Val Asp Pro Val Pro
 290 295 300
 Ser Tyr Lys Gln Val Lys Gln Leu Lys Tyr Val Gly Met Val Leu Asn
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 Glu Ala Leu Arg Leu Trp Pro Thr Ala Pro Ala Phe Ser Leu Tyr Ala
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 Lys Glu Asp Thr Val Leu Gly Gly Glu Tyr Pro Leu Glu Lys Gly Asp
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 Glu Leu Met Val Leu Ile Pro Gln Leu His Arg Asp Lys Thr Ile Trp
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 Gly Asp Asp Val Glu Glu Phe Arg Pro Glu Arg Phe Glu Asn Pro Ser
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 Glu Gln Ser Ala Lys Lys Val Arg Lys Lys Ala Glu Asn Ala His Asn
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 Thr Ala Arg Asp Leu Ala Asp Ile Ala Met Ser Lys Gly Phe Ala Pro
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 Gln Val Ala Thr Leu Asp Ser His Ala Gly Asn Leu Pro Arg Glu Gly
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 Asp Phe Glu Gly Thr Tyr Glu Glu Trp Arg Glu His Met Trp Ser Asp
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Leu Pro Lys Glu Ala Ser Tyr 690	Gln Glu Gly Asp 695	His Leu Gly Val Ile 700
Pro Arg Asn Tyr Glu Gly Ile 705	Val Asn Arg Val Thr 710	Ala Arg Phe Gly 715
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Ala His Leu Pro Leu Ala Lys 740	Thr Val Ser Val Glu Glu 745	Leu Leu Gln 750
Tyr Val Glu Leu Gln Asp 755	Pro Val Thr Arg Thr 760	Gln Leu Arg Ala Met 765
Ala Ala Lys Thr Val Cys 770	Pro His Lys Val Glu 775	Leu Leu Glu Ala Leu 780
Leu Glu Lys Gln Ala Tyr 785	Lys Glu Gln Val Leu 790	Ala Lys Arg Leu Thr 795
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Glu Phe Ile Ala Leu Leu 820	Pro Ser Ile Arg Pro 825	Arg Tyr Tyr Ser Ile 830
Ser Ser Ser Pro Arg Val 835	Asp Glu Lys Gln Ala 840	Ser Ile Thr Val Ser 845
Val Val Ser Gly Glu Ala 850	Trp Ser Gly Tyr Gly 855	Glu Tyr Lys Gly Ile 860
Ala Ser Asn Tyr Leu Ala 865	Glu Leu Gln Glu Gly 870	Asp Thr Ile Thr Cys 875
Phe Ile Ser Thr Pro Gln 885	Ser Glu Phe Thr Leu 890	Pro Lys Asp Pro Glu 895
Thr Pro Leu Ile Met Val 900	Gly Pro Gly Thr Gly 905	Val Ala Pro Phe Arg 910
Gly Phe Val Gln Ala Arg 915	Lys Gln Leu Lys Glu 920	Gln Gln Ser Leu 925
Gly Glu Ala His Leu Tyr 930	Phe Gly Cys Arg Ser 935	Pro His Glu Asp Tyr 940
Leu Tyr Gln Glu Glu Leu 945	Glu Asn Ala Gln Ser 950	Glu Gly Ile Ile Thr 955
Leu His Thr Ala Phe Ser 965	Arg Met Pro Asn Gln 970	Pro Lys Thr Tyr Val 975
Gln His Val Met Glu Gln 980	Asp Gly Lys Lys Leu 985	Ile Glu Leu Leu Asp 990
Gln Gly Ala His Phe Tyr 995	Ile Cys Gly Asp Gly 1000	Ser Gln Met Ala Pro 1005
Ala Val Glu Ala Thr Leu 1010	Met Lys Ser Tyr Ala 1015	Asp Val His Gln 1020
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 Ala Asp Glu Leu Gly Glu Ile Phe Lys Phe Glu Ala Pro Gly Arg Val
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 Thr Arg Tyr Leu Ser Ser Gln Arg Leu Ile Lys Glu Ala Cys Asp Glu
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 Ser Arg Phe Asp Lys Asn Leu Ser Gln Ala Leu Lys Phe Val Arg Asp
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 85 90 95
 aaa aaa gcg cat aat atc tta ctt cca agc ttc agt cag cag gca atg 336
 Lys Lys Ala His Asn Ile Leu Leu Pro Ser Phe Ser Gln Gln Ala Met
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 aaa ggc tat cat gcg atg atg gtc gat atc gcc gtg cag ctt gtt caa 384
 Lys Gly Tyr His Ala Met Met Val Asp Ile Ala Val Gln Leu Val Gln
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 aag tgg gag cgt cta aat gca gat gag cat att gaa gta ccg gaa gac 432
 Lys Trp Glu Arg Leu Asn Ala Asp Glu His Ile Glu Val Pro Glu Asp
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 atg aca cgt tta acg ctt gat aca att ggt ctt tgc ggc ttt aac tat 480
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 Arg Phe Asn Ser Phe Tyr Arg Asp Gln Pro His Pro Phe Ile Thr Ser
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 Pro Asp Asp Pro Ala Tyr Asp Glu Asn Lys Arg Gln Phe Gln Glu Asp
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 Ile Lys Val Met Asn Asp Leu Val Asp Lys Ile Ile Ala Asp Arg Lys
 210 215 220
 gca agc ggt gaa caa agc gat gat tta tta acg cat atg cta aac gga 720
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ggc gtt cgc tac tcc gta ttt gga tgc ggc gat aaa aac tgg gct act				1728
Gly Val Arg Tyr	Ser Val Phe Gly Cys	Gly Asp Lys Asn Trp	Ala Thr	
565	570	575		
acg tat caa aaa gtg cct gct ttt atc gat gaa acg ctt gcc gct aaa				1776
Thr Tyr Gln	Lys Val Pro Ala Phe	Ile Asp Glu Thr Leu	Ala Ala Lys	
580	585	590		
ggg gca gaa aac atc gct gac cgc ggt gaa gca gat gca agc gac gac				1824
Gly Ala Glu Asn Ile Ala Asp	Arg Gly Glu Ala Asp	Ala Ser Asp Asp		
595	600	605		
ttt gaa ggc aca tat gaa gaa tgg cgt gaa cat atg tgg agt gac gta				1872
Phe Glu Gly Thr Tyr Glu Glu Trp Arg	Glu His Met	Trp Ser Asp Val		
610	615	620		
gca gcc tac ttt aac ctc gac att gaa aac agt gaa gat aat aaa tct				1920
Ala Ala Tyr Phe Asn Leu Asp	Ile Glu Asn Ser	Glu Asp Asn Lys Ser		
625	630	635	640	
act ctt tca ctt caa ttt gtc gac agc gcc gcg gat atg ccg ctt gcg				1968
Thr Leu Ser Leu Gln Phe Val Asp	Ser Ala Ala Asp Met	Pro Leu Ala		
645	650	655		
aaa atg cac ggt gcg ttt tca acg aac gtc gta gca agc aaa gaa ctt				2016
Lys Met His Gly Ala Phe Ser Thr Asn Val Val Ala Ser	Lys Glu Leu			
660	665	670		
caa cag cca ggc agt gca cga agc acg cga cat ctt gaa att gaa ctt				2064
Gln Gln Pro Gly Ser Ala Arg Ser Thr Arg His Leu	Glu Ile Glu Leu			
675	680	685		
cca aaa gaa gct tct tat caa gaa gga gat cat tta ggt gtt att cct				2112
Pro Lys Glu Ala Ser Tyr Gln Glu Gly Asp His Leu Gly Val Ile Pro				
690	695	700		
cgc aac tat gaa gga ata gta aac cgt gta aca gca agg ttc ggc cta				2160
Arg Asn Tyr Glu Gly Ile Val Asn Arg Val Thr Ala Arg Phe Gly Leu				
705	710	715	720	
gat gca tca cag caa atc cgt ctg gaa gca gaa gaa gaa aaa tta gct				2208
Asp Ala Ser Gln Gln Ile Arg Leu Glu Ala Glu Glu Glu Lys Leu Ala				
725	730	735		
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His Leu Pro Leu Ala Lys Thr Val Ser Val Glu Glu Leu Leu Gln Tyr				
740	745	750		
gtg gag ctt caa gat cct gtt acg cgc acg cag ctt cgc gca atg gct				2304
Val Glu Leu Gln Asp Pro Val Thr Arg Thr Gln Leu Arg Ala Met Ala				
755	760	765		
gct aaa acg gtc tgc ccg ccg cat aaa gta gag ctt gaa gcc ttg ctt				2352
Ala Lys Thr Val Cys Pro Pro His Lys Val Glu Leu Glu Ala Leu Leu				
770	775	780		
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Glu Lys Gln Ala Tyr Lys Glu Gln Val Leu Ala Lys Arg Leu Thr Met				
785	790	795	800	
ctt gaa ctg ctt gaa aaa tac ccg gcg tgt gaa atg aaa ttc agc gaa				2448
Leu Glu Leu Leu Glu Lys Tyr Pro Ala Cys Glu Met Lys Phe Ser Glu				
805	810	815		
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Phe Ile Ala Leu Leu Pro Ser Ile Arg Pro Arg Tyr Tyr Ser Ile Ser				
820	825	830		
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Ser Ser Pro Arg Val Asp Glu Lys Gln Ala Ser Ile Thr Val Ser Val				
835	840	845		
gtc tca gga gaa gcg tgg agc gga tat gga gaa tat aaa gga att gcg				2592
Val Ser Gly Glu Ala Trp Ser Gly Tyr Gly Glu Tyr Lys Gly Ile Ala				

850 855 860

tcg aac tat ctt gcc gag ctg caa gaa gga gat acg att acg tgc ttt 2640
 Ser Asn Tyr Leu Ala Glu Leu Gln Glu Gly Asp Thr Ile Thr Cys Phe
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 Ile Ser Thr Pro Gln Ser Glu Phe Thr Leu Pro Lys Asp Pro Glu Thr
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gaa gca cat tta tac ttc ggc tgc cgt tca cct cat gaa gac tat ctg 2832
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 Tyr Gln Glu Glu Leu Glu Asn Ala Gln Ser Glu Gly Ile Ile Thr Leu
 945 950 955 960

cat acc gct ttt tct cgc atg cca aat cag ccg aaa aca tac gtt cag 2928
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 965 970 975

cac gta atg gaa caa gac ggc aag aaa ttg att gaa ctt ctt gat caa 2976
 His Val Met Glu Gln Asp Gly Lys Lys Leu Ile Glu Leu Leu Asp Gln
 980 985 990

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 995 1000 1005

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 35 40 45

Thr Arg Tyr Leu Ser Ser Gln Arg Leu Ile Lys Glu Ala Cys Asp Glu
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Ser Arg Phe Asp Lys Asn Leu Ser Gln Ala Leu Lys Phe Val Arg Asp
 65 70 75 80

Phe Ala Gly Asp Gly Leu Ala Thr Ser Trp Thr His Glu Lys Asn Trp
 85 90 95
 Lys Lys Ala His Asn Ile Leu Leu Pro Ser Phe Ser Gln Gln Ala Met
 100 105 110
 Lys Gly Tyr His Ala Met Met Val Asp Ile Ala Val Gln Leu Val Gln
 115 120 125
 Lys Trp Glu Arg Leu Asn Ala Asp Glu His Ile Glu Val Pro Glu Asp
 130 135 140
 Met Thr Arg Leu Thr Leu Asp Thr Ile Gly Leu Cys Gly Phe Asn Tyr
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 Arg Phe Asn Ser Phe Tyr Arg Asp Gln Pro His Pro Phe Ile Thr Ser
 165 170 175
 Met Val Arg Ala Leu Asp Glu Ala Met Asn Lys Leu Gln Arg Ala Asn
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 195 200 205
 Ile Lys Val Met Asn Asp Leu Val Asp Lys Ile Ile Ala Asp Arg Lys
 210 215 220
 Ala Ser Gly Glu Gln Ser Asp Asp Leu Leu Thr His Met Leu Asn Gly
 225 230 235 240
 Lys Asp Pro Glu Thr Gly Glu Pro Leu Asp Asp Glu Asn Ile Arg Tyr
 245 250 255
 Gln Ile Ile Thr Phe Leu Ile Ala Gly His Glu Thr Thr Ser Gly Leu
 260 265 270
 Leu Ser Phe Ala Leu Tyr Phe Leu Val Lys Asn Pro His Val Leu Gln
 275 280 285
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 Tyr Lys Gln Val Lys Gln Leu Lys Tyr Val Gly Met Val Leu Asn Glu
 305 310 315 320
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 325 330 335
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 340 345 350
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 355 360 365
 Asp Asp Val Glu Glu Phe Arg Pro Glu Arg Phe Glu Asn Pro Ser Ala
 370 375 380
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 385 390 395 400
 Ile Gly Gln Gln Phe Ala Leu His Glu Ala Thr Leu Val Leu Gly Met
 405 410 415
 Met Leu Lys His Phe Asp Phe Glu Asp His Thr Asn Tyr Glu Leu Asp
 420 425 430
 Ile Lys Glu Thr Leu Thr Leu Lys Pro Glu Gly Phe Val Val Lys Ala
 435 440 445
 Lys Ser Lys Lys Ile Pro Leu Gly Gly Ile Pro Ser Pro Ser Thr Glu
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 Gln Ser Ala Lys Lys Val Arg Lys Lys Ala Glu Asn Ala His Asn Thr
 465 470 475 480
 Pro Leu Leu Val Leu Tyr Gly Ser Asn Met Gly Thr Ala Glu Gly Thr

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Val Leu Ile Val Thr Ala Ser Tyr Asn Gly His Pro Pro Asp Asn Ala 530 535 540		
Lys Gln Phe Val Asp Trp Leu Asp Gln Ala Ser Ala Asp Glu Val Lys 545 550 555 560		
Gly Val Arg Tyr Ser Val Phe Gly Cys Gly Asp Lys Asn Trp Ala Thr 565 570 575		
Thr Tyr Gln Lys Val Pro Ala Phe Ile Asp Glu Thr Leu Ala Ala Lys 580 585 590		
Gly Ala Glu Asn Ile Ala Asp Arg Gly Glu Ala Asp Ala Ser Asp Asp 595 600 605		
Phe Glu Gly Thr Tyr Glu Glu Trp Arg Glu His Met Trp Ser Asp Val 610 615 620		
Ala Ala Tyr Phe Asn Leu Asp Ile Glu Asn Ser Glu Asp Asn Lys Ser 625 630 635 640		
Thr Leu Ser Leu Gln Phe Val Asp Ser Ala Ala Asp Met Pro Leu Ala 645 650 655		
Lys Met His Gly Ala Phe Ser Thr Asn Val Val Ala Ser Lys Glu Leu 660 665 670		
Gln Gln Pro Gly Ser Ala Arg Ser Thr Arg His Leu Glu Ile Glu Leu 675 680 685		
Pro Lys Glu Ala Ser Tyr Gln Glu Gly Asp His Leu Gly Val Ile Pro 690 695 700		
Arg Asn Tyr Glu Gly Ile Val Asn Arg Val Thr Ala Arg Phe Gly Leu 705 710 715 720		
Asp Ala Ser Gln Gln Ile Arg Leu Glu Ala Glu Glu Glu Lys Leu Ala 725 730 735		
His Leu Pro Leu Ala Lys Thr Val Ser Val Glu Glu Leu Leu Gln Tyr 740 745 750		
Val Glu Leu Gln Asp Pro Val Thr Arg Thr Gln Leu Arg Ala Met Ala 755 760 765		
Ala Lys Thr Val Cys Pro Pro His Lys Val Glu Leu Glu Ala Leu Leu 770 775 780		
Glu Lys Gln Ala Tyr Lys Glu Gln Val Leu Ala Lys Arg Leu Thr Met 785 790 795 800		
Leu Glu Leu Leu Glu Lys Tyr Pro Ala Cys Glu Met Lys Phe Ser Glu 805 810 815		
Phe Ile Ala Leu Leu Pro Ser Ile Arg Pro Arg Tyr Tyr Ser Ile Ser 820 825 830		
Ser Ser Pro Arg Val Asp Glu Lys Gln Ala Ser Ile Thr Val Ser Val 835 840 845		
Val Ser Gly Glu Ala Trp Ser Gly Tyr Gly Glu Tyr Lys Gly Ile Ala 850 855 860		
Ser Asn Tyr Leu Ala Glu Leu Gln Glu Gly Asp Thr Ile Thr Cys Phe 865 870 875 880		
Ile Ser Thr Pro Gln Ser Glu Phe Thr Leu Pro Lys Asp Pro Glu Thr 885 890 895		

Pro Leu Ile Met Val Gly Pro Gly Thr Gly Val Ala Pro Phe Arg Gly
 900 905 910

Phe Val Gln Ala Arg Lys Gln Leu Lys Glu Gln Gly Gln Ser Leu Gly
 915 920 925

Glu Ala His Leu Tyr Phe Gly Cys Arg Ser Pro His Glu Asp Tyr Leu
 930 935 940

Tyr Gln Glu Glu Leu Glu Asn Ala Gln Ser Glu Gly Ile Ile Thr Leu
 945 950 955 960

His Thr Ala Phe Ser Arg Met Pro Asn Gln Pro Lys Thr Tyr Val Gln
 965 970 975

His Val Met Glu Gln Asp Gly Lys Lys Leu Ile Glu Leu Leu Asp Gln
 980 985 990

Gly Ala His Phe Tyr Ile Cys Gly Asp Gly Ser Gln Met Ala Pro Ala
 995 1000 1005

Val Glu Ala Thr Leu Met Lys Ser Tyr Ala Asp Val His Gln Val
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 20 25 30

gcg gat gaa tta gga gaa atc ttt aaa ttc gag gcg cct ggt cgt gta 144
 Ala Asp Glu Leu Gly Glu Ile Phe Lys Phe Glu Ala Pro Gly Arg Val
 35 40 45

acg cgc tac tta tca agt cag cgt cta att aaa gaa gca tgc gat gaa 192
 Thr Arg Tyr Leu Ser Ser Gln Arg Leu Ile Lys Glu Ala Cys Asp Glu
 50 55 60

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 Ser Arg Phe Asp Lys Asn Leu Ser Gln Gly Leu Lys Phe Val Arg Asp
 65 70 75 80

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 85 90 95

aaa aaa gcg cat aat atc tta ctt cca agc ttc agt cag cag gca atg 336
 Lys Lys Ala His Asn Ile Leu Leu Pro Ser Phe Ser Gln Gln Ala Met
 100 105 110

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Lys Gly Tyr His Ala Met Met Val Asp Ile Ala Val Gln Leu Val Gln	
115 120 125	
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Lys Trp Glu Arg Leu Asn Ala Asp Glu His Ile Glu Val Pro Glu Asp	
130 135 140	
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Met Thr Arg Leu Thr Leu Asp Thr Ile Gly Leu Cys Gly Phe Asn Tyr	
145 150 155 160	
cgc ttt aac agc ttt tac cga gat cag cct cat cca ttt att aca agt	528
Arg Phe Asn Ser Phe Tyr Arg Asp Gln Pro His Pro Phe Ile Thr Ser	
165 170 175	
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Pro Asp Asp Pro Ala Tyr Asp Glu Asn Lys Arg Gln Phe Gln Glu Asp	
195 200 205	
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Ile Lys Val Met Asn Asp Leu Val Asp Lys Ile Ile Ala Asp Arg Lys	
210 215 220	
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Lys Asp Pro Glu Thr Gly Glu Pro Leu Asp Asp Glu Asn Ile Arg Tyr	
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Gln Ile Ile Thr Phe Leu Ile Ala Gly His Glu Thr Thr Ser Gly Leu	
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Leu Ser Phe Ala Leu Tyr Phe Leu Val Lys Asn Pro His Val Leu Gln	
275 280 285	
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Tyr Lys Gln Val Lys Gln Leu Lys Tyr Val Gly Met Val Leu Asn Glu	
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Tyr Gln Glu Glu Leu Glu Asn Ala Gln Ser Glu Gly Ile Ile Thr Leu	
945 950 955 960	
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His Val Met Glu Gln Asp Gly Lys Lys Leu Ile Glu Leu Leu Asp Gln	
980 985 990	
gga gcg cac ttc tat att tgc gga gac gga agc caa atg gca cct gcc	3024
Gly Ala His Phe Tyr Ile Cys Gly Asp Gly Ser Gln Met Ala Pro Ala	
995 1000 1005	
gtt gaa gca acg ctt atg aaa agc tat gct gac gtt cac caa gtg	3069
Val Glu Ala Thr Leu Met Lys Ser Tyr Ala Asp Val His Gln Val	
1010 1015 1020	
agt gaa gca gac gct cgc tta tgg ctg cag cag cta gaa gaa aaa	3114

Ser Glu Ala Asp Ala Arg Leu Trp Leu Gln Gln Leu Glu Glu Lys
 1025 1030 1035

ggc cga tac gca aaa gac gtg tgg gct ggg taa
 Gly Arg Tyr Ala Lys Asp Val Trp Ala Gly
 1040 1045

3147

<210> 8
 <211> 1048
 <212> PRT
 <213> Artificial sequence

<220>
 <223> Cytochrome P450BM-3 mutant

<400> 8

Thr Ile Lys Glu Met Pro Gln Pro Lys Thr Phe Gly Glu Leu Lys Asn
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 Leu Pro Leu Leu Asn Thr Asp Lys Pro Val Gln Ala Leu Met Lys Ile
 20 25 30
 Ala Asp Glu Leu Gly Glu Ile Phe Lys Phe Glu Ala Pro Gly Arg Val
 35 40 45
 Thr Arg Tyr Leu Ser Ser Gln Arg Leu Ile Lys Glu Ala Cys Asp Glu
 50 55 60
 Ser Arg Phe Asp Lys Asn Leu Ser Gln Gly Leu Lys Phe Val Arg Asp
 65 70 75 80
 Phe Ala Gly Asp Gly Leu Val Thr Ser Trp Thr His Glu Lys Asn Trp
 85 90 95
 Lys Lys Ala His Asn Ile Leu Leu Pro Ser Phe Ser Gln Gln Ala Met
 100 105 110
 Lys Gly Tyr His Ala Met Met Val Asp Ile Ala Val Gln Leu Val Gln
 115 120 125
 Lys Trp Glu Arg Leu Asn Ala Asp Glu His Ile Glu Val Pro Glu Asp
 130 135 140
 Met Thr Arg Leu Thr Leu Asp Thr Ile Gly Leu Cys Gly Phe Asn Tyr
 145 150 155 160
 Arg Phe Asn Ser Phe Tyr Arg Asp Gln Pro His Pro Phe Ile Thr Ser
 165 170 175
 Met Val Arg Ala Leu Asp Glu Ala Met Asn Lys Gln Gln Arg Ala Asn
 180 185 190
 Pro Asp Asp Pro Ala Tyr Asp Glu Asn Lys Arg Gln Phe Gln Glu Asp
 195 200 205
 Ile Lys Val Met Asn Asp Leu Val Asp Lys Ile Ile Ala Asp Arg Lys
 210 215 220
 Ala Ser Gly Glu Gln Ser Asp Asp Leu Leu Thr His Met Leu Asn Gly
 225 230 235 240
 Lys Asp Pro Glu Thr Gly Glu Pro Leu Asp Asp Glu Asn Ile Arg Tyr
 245 250 255
 Gln Ile Ile Thr Phe Leu Ile Ala Gly His Glu Thr Thr Ser Gly Leu
 260 265 270
 Leu Ser Phe Ala Leu Tyr Phe Leu Val Lys Asn Pro His Val Leu Gln
 275 280 285
 Lys Ala Ala Glu Glu Ala Ala Arg Val Leu Val Asp Pro Val Pro Ser
 290 295 300

Tyr Lys Gln Val Lys Gln Leu Lys Tyr Val Gly Met Val Leu Asn Glu
 305 310 315 320
 Ala Leu Arg Leu Trp Pro Thr Ala Pro Ala Phe Ser Leu Tyr Ala Lys
 325 330 335
 Glu Asp Thr Val Leu Gly Gly Glu Tyr Pro Leu Glu Lys Gly Asp Glu
 340 345 350
 Leu Met Val Leu Ile Pro Gln Leu His Arg Asp Lys Thr Ile Trp Gly
 355 360 365
 Asp Asp Val Glu Glu Phe Arg Pro Glu Arg Phe Glu Asn Pro Ser Ala
 370 375 380
 Ile Pro Gln His Ala Phe Lys Pro Phe Gly Asn Gly Gln Arg Ala Cys
 385 390 395 400
 Ile Gly Gln Gln Phe Ala Leu His Glu Ala Thr Leu Val Leu Gly Met
 405 410 415
 Met Leu Lys His Phe Asp Phe Glu Asp His Thr Asn Tyr Glu Leu Asp
 420 425 430
 Ile Lys Glu Thr Leu Thr Leu Lys Pro Glu Gly Phe Val Val Lys Ala
 435 440 445
 Lys Ser Lys Lys Ile Pro Leu Gly Gly Ile Pro Ser Pro Ser Thr Glu
 450 455 460
 Gln Ser Ala Lys Lys Val Arg Lys Lys Ala Glu Asn Ala His Asn Thr
 465 470 475 480
 Pro Leu Leu Val Leu Tyr Gly Ser Asn Met Gly Thr Ala Glu Gly Thr
 485 490 495
 Ala Arg Asp Leu Ala Asp Ile Ala Met Ser Lys Gly Phe Ala Pro Gln
 500 505 510
 Val Ala Thr Leu Asp Ser His Ala Gly Asn Leu Pro Arg Glu Gly Ala
 515 520 525
 Val Leu Ile Val Thr Ala Ser Tyr Asn Gly His Pro Pro Asp Asn Ala
 530 535 540
 Lys Gln Phe Val Asp Trp Leu Asp Gln Ala Ser Ala Asp Glu Val Lys
 545 550 555 560
 Gly Val Arg Tyr Ser Val Phe Gly Cys Gly Asp Lys Asn Trp Ala Thr
 565 570 575
 Thr Tyr Gln Lys Val Pro Ala Phe Ile Asp Glu Thr Leu Ala Ala Lys
 580 585 590
 Gly Ala Glu Asn Ile Ala Asp Arg Gly Glu Ala Asp Ala Ser Asp Asp
 595 600 605
 Phe Glu Gly Thr Tyr Glu Glu Trp Arg Glu His Met Trp Ser Asp Val
 610 615 620
 Ala Ala Tyr Phe Asn Leu Asp Ile Glu Asn Ser Glu Asp Asn Lys Ser
 625 630 635 640
 Thr Leu Ser Leu Gln Phe Val Asp Ser Ala Ala Asp Met Pro Leu Ala
 645 650 655
 Lys Met His Gly Ala Phe Ser Thr Asn Val Val Ala Ser Lys Glu Leu
 660 665 670
 Gln Gln Pro Gly Ser Ala Arg Ser Thr Arg His Leu Glu Ile Glu Leu
 675 680 685
 Pro Lys Glu Ala Ser Tyr Gln Glu Gly Asp His Leu Gly Val Ile Pro
 690 695 700
 Arg Asn Tyr Glu Gly Ile Val Asn Arg Val Thr Ala Arg Phe Gly Leu

705 710 715 720
 Asp Ala Ser Gln Gln Ile Arg Leu Glu Ala Glu Glu Glu Lys Leu Ala
 725 730 735
 His Leu Pro Leu Ala Lys Thr Val Ser Val Glu Glu Leu Leu Gln Tyr
 740 745 750
 Val Glu Leu Gln Asp Pro Val Thr Arg Thr Gln Leu Arg Ala Met Ala
 755 760 765
 Ala Lys Thr Val Cys Pro Pro His Lys Val Glu Leu Glu Ala Leu Leu
 770 775 780
 Glu Lys Gln Ala Tyr Lys Glu Gln Val Leu Ala Lys Arg Leu Thr Met
 785 790 795 800
 Leu Glu Leu Leu Glu Lys Tyr Pro Ala Cys Glu Met Lys Phe Ser Glu
 805 810 815
 Phe Ile Ala Leu Leu Pro Ser Ile Arg Pro Arg Tyr Tyr Ser Ile Ser
 820 825 830
 Ser Ser Pro Arg Val Asp Glu Lys Gln Ala Ser Ile Thr Val Ser Val
 835 840 845
 Val Ser Gly Glu Ala Trp Ser Gly Tyr Gly Glu Tyr Lys Gly Ile Ala
 850 855 860
 Ser Asn Tyr Leu Ala Glu Leu Gln Glu Gly Asp Thr Ile Thr Cys Phe
 865 870 875 880
 Ile Ser Thr Pro Gln Ser Glu Phe Thr Leu Pro Lys Asp Pro Glu Thr
 885 890 895
 Pro Leu Ile Met Val Gly Pro Gly Thr Gly Val Ala Pro Phe Arg Gly
 900 905 910
 Phe Val Gln Ala Arg Lys Gln Leu Lys Glu Gln Gly Gln Ser Leu Gly
 915 920 925
 Glu Ala His Leu Tyr Phe Gly Cys Arg Ser Pro His Glu Asp Tyr Leu
 930 935 940
 Tyr Gln Glu Glu Leu Glu Asn Ala Gln Ser Glu Gly Ile Ile Thr Leu
 945 950 955 960
 His Thr Ala Phe Ser Arg Met Pro Asn Gln Pro Lys Thr Tyr Val Gln
 965 970 975
 His Val Met Glu Gln Asp Gly Lys Lys Leu Ile Glu Leu Leu Asp Gln
 980 985 990
 Gly Ala His Phe Tyr Ile Cys Gly Asp Gly Ser Gln Met Ala Pro Ala
 995 1000 1005
 Val Glu Ala Thr Leu Met Lys Ser Tyr Ala Asp Val His Gln Val
 1010 1015 1020
 Ser Glu Ala Asp Ala Arg Leu Trp Leu Gln Gln Leu Glu Glu Lys
 1025 1030 1035
 Gly Arg Tyr Ala Lys Asp Val Trp Ala Gly
 1040 1045

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 <211> 1032
 <212> DNA
 <213> Nocardia corallina

<220>
 <221> CDS
 <222> (1)..(1032)

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cgg aca ttc acc tgg ttc acg ccc gcc agg cga aag ccg acg gag tac		96
Arg Thr Phe Thr Trp Phe Thr Pro Ala Arg Arg Lys Pro Thr Glu Tyr		
20 25 30		
gag ctc tac acc gtg ggt caa cag tcc act ccg gac gag tgg ctg cat		144
Glu Leu Tyr Thr Val Gly Gln Gln Ser Thr Pro Asp Glu Trp Leu His		
35 40 45		
gtg gac tgg ccg ctg cgc ttc gac gac ggc cgc gcc ccg tgg gag gag		192
Val Asp Trp Pro Leu Arg Phe Asp Asp Gly Arg Ala Pro Trp Glu Glu		
50 55 60		
gag tcg agt gcg gta cgg acc tcg gag tgg tgc gct tac cgc gac cca		240
Glu Ser Ser Ala Val Arg Thr Ser Glu Trp Ser Ala Tyr Arg Asp Pro		
65 70 75 80		
cac caa ctg tgg cag cgt ccc tac gtc agc acg tgc aac cag gac cag		288
His Gln Leu Trp Gln Arg Pro Tyr Val Ser Thr Cys Asn Gln Asp Gln		
85 90 95		
cag gcc ctc gcg cgg ctg gtc ccc gtc ctg acc atg ggg tgc gcg gcg		336
Gln Ala Leu Ala Arg Leu Val Pro Val Leu Thr Met Gly Ser Ala Ala		
100 105 110		
atc acg ccc atc tgg tgc cag aag atc ctc gcc agg tcc tac gcc gcc		384
Ile Thr Pro Ile Trp Ser Gln Lys Ile Leu Ala Arg Ser Tyr Ala Ala		
115 120 125		
tgg cca ttc gtc gag tac ggg ctc ttc ctg agc ctg gcc tac gcc gtg		432
Trp Pro Phe Val Glu Tyr Gly Leu Phe Leu Ser Leu Ala Tyr Ala Val		
130 135 140		
cgc cag gcc atg tcc gac acg gtc cag ttc agc gtg gtg ttc cag gcc		480
Arg Gln Ala Met Ser Asp Thr Val Gln Phe Ser Val Val Phe Gln Ala		
145 150 155 160		
gtg gac cgc atg cgg ctg ctc cag gac atc gtc cac cac ctg gac cac		528
Val Asp Arg Met Arg Leu Leu Gln Asp Ile Val His His Leu Asp His		
165 170 175		
ctg cag gag tgc ccg gaa ttc agc gac gcc ggg gcc cgc gag gcc tgg		576
Leu Gln Glu Ser Pro Glu Phe Ser Asp Ala Gly Ala Arg Glu Ala Trp		
180 185 190		
atg tcc gac tcc acc ctg gtc ccg atc cgg gaa gtg atc gag cgc atc		624
Met Ser Asp Ser Thr Leu Val Pro Ile Arg Glu Val Ile Glu Arg Ile		
195 200 205		
gcc gcc agc cag gac tgg gtg gag atc ctg gtc gcc ggc acg ctc gtc		672
Ala Ala Ser Gln Asp Trp Val Glu Ile Leu Val Ala Gly Thr Leu Val		
210 215 220		
ttc gag cct ctg gtc ggc cac ctg gcg aag gcc gag ttg ttc agc cgc		720
Phe Glu Pro Leu Val Gly His Leu Ala Lys Ala Glu Leu Phe Ser Arg		
225 230 235 240		
cgt gcg cca atg ttc ggg gac ggg acc acg ccg gcg gtg ctg gcg tgc		768
Arg Ala Pro Met Phe Gly Asp Gly Thr Thr Pro Ala Val Leu Ala Ser		
245 250 255		
gcc ctg ctg gac agc ggc agg cac ctc gaa tgc gtc cag gcg ctc gtc		816
Ala Leu Leu Asp Ser Gly Arg His Leu Glu Ser Val Gln Ala Leu Val		
260 265 270		
cgc ctc gtc tgc caa gac ccc gtc cat ggc gac cag aac cag gcg act		864
Arg Leu Val Cys Gln Asp Pro Val His Gly Asp Gln Asn Gln Ala Thr		
275 280 285		
gtg cgg cgg tgg atc gag gaa tgg cag ccg cgg tgc aag gcg gcg gcc		912
Val Arg Arg Trp Ile Glu Glu Trp Gln Pro Arg Cys Lys Ala Ala Ala		
290 295 300		

cag tcc ttc ctg ccg acg ttc tcc gac tgc ggc atc gac gcc aag gaa 960
 Gln Ser Phe Leu Pro Thr Phe Ser Asp Cys Gly Ile Asp Ala Lys Glu
 305 310 315 320

agc gcc aac gcg ctg tcc cgg gcg ctg gcg aac cag cgg gcc gcc gtc 1008
 Ser Ala Asn Ala Leu Ser Arg Ala Leu Ala Asn Gln Arg Ala Ala Val
 325 330 335

gag ggc gcc ggc atc acg gca tga 1032
 Glu Gly Ala Gly Ile Thr Ala
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<210> 10
 <211> 343
 <212> PRT
 <213> Nocardia corallina

<400> 10

Met Thr Thr Glu Ala Thr Val Ala Arg Pro Val Glu Leu Glu Gly His
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Arg Thr Phe Thr Trp Phe Thr Pro Ala Arg Arg Lys Pro Thr Glu Tyr
 20 25 30

Glu Leu Tyr Thr Val Gly Gln Gln Ser Thr Pro Asp Glu Trp Leu His
 35 40 45

Val Asp Trp Pro Leu Arg Phe Asp Asp Gly Arg Ala Pro Trp Glu Glu
 50 55 60

Glu Ser Ser Ala Val Arg Thr Ser Glu Trp Ser Ala Tyr Arg Asp Pro
 65 70 75 80

His Gln Leu Trp Gln Arg Pro Tyr Val Ser Thr Cys Asn Gln Asp Gln
 85 90 95

Gln Ala Leu Ala Arg Leu Val Pro Val Leu Thr Met Gly Ser Ala Ala
 100 105 110

Ile Thr Pro Ile Trp Ser Gln Lys Ile Leu Ala Arg Ser Tyr Ala Ala
 115 120 125

Trp Pro Phe Val Glu Tyr Gly Leu Phe Leu Ser Leu Ala Tyr Ala Val
 130 135 140

Arg Gln Ala Met Ser Asp Thr Val Gln Phe Ser Val Val Phe Gln Ala
 145 150 155 160

Val Asp Arg Met Arg Leu Leu Gln Asp Ile Val His His Leu Asp His
 165 170 175

Leu Gln Glu Ser Pro Glu Phe Ser Asp Ala Gly Ala Arg Glu Ala Trp
 180 185 190

Met Ser Asp Ser Thr Leu Val Pro Ile Arg Glu Val Ile Glu Arg Ile
 195 200 205

Ala Ala Ser Gln Asp Trp Val Glu Ile Leu Val Ala Gly Thr Leu Val
 210 215 220

Phe Glu Pro Leu Val Gly His Leu Ala Lys Ala Glu Leu Phe Ser Arg
 225 230 235 240

Arg Ala Pro Met Phe Gly Asp Gly Thr Thr Pro Ala Val Leu Ala Ser
 245 250 255

Ala Leu Leu Asp Ser Gly Arg His Leu Glu Ser Val Gln Ala Leu Val
 260 265 270

Arg Leu Val Cys Gln Asp Pro Val His Gly Asp Gln Asn Gln Ala Thr
 275 280 285

Val Arg Arg Trp Ile Glu Glu Trp Gln Pro Arg Cys Lys Ala Ala Ala

290 295 300

Gln Ser Phe Leu Pro Thr Phe Ser Asp Cys Gly Ile Asp Ala Lys Glu
305 310 315 320

Ser Ala Asn Ala Leu Ser Arg Ala Leu Ala Asn Gln Arg Ala Ala Val
325 330 335

Glu Gly Ala Gly Ile Thr Ala
340

<210> 11
<211> 1506
<212> DNA
<213> *Nocardia corallina*

<220>
<221> CDS
<222> (1)..(1506)

<400> 11

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tgg gac ttc acc tcc gtc gag cgg cgc ccc aag ttc gag acg aag tac	96
Trp Asp Phe Thr Ser Val Glu Arg Arg Pro Lys Phe Glu Thr Lys Tyr	
20 25 30	
aag atg ccc aag aag ggc aag gac ccg ttc cgc gtc ctg atc cgt gac	144
Lys Met Pro Lys Lys Gly Lys Asp Pro Phe Arg Val Leu Ile Arg Asp	
35 40 45	
tac atg aag atg gaa gcg gag aag gac gac cgg acc cat ggc ttc ctc	192
Tyr Met Lys Met Glu Ala Glu Lys Asp Asp Arg Thr His Gly Phe Leu	
50 55 60	
gac ggc gcc gtg cgg acg cgt gag gcc acc agg att gag ccg cgg ttc	240
Asp Gly Ala Val Arg Thr Arg Glu Ala Thr Arg Ile Glu Pro Arg Phe	
65 70 75 80	
gct gag gcc atg aag atc atg gtg ccg cag ctg acc aac gcc gag tac	288
Ala Glu Ala Met Lys Ile Met Val Pro Gln Leu Thr Asn Ala Glu Tyr	
85 90 95	
cag gcg gtg gcg ggc tgc gga atg atc atc tcg gcc gtc gag aac cag	336
Gln Ala Val Ala Gly Cys Gly Met Ile Ile Ser Ala Val Glu Asn Gln	
100 105 110	
gag ctc cgt cag ggc tac gcc gct cag atg ctc gat gag gtg cgg cac	384
Glu Leu Arg Gln Gly Tyr Ala Ala Gln Met Leu Asp Glu Val Arg His	
115 120 125	
gcg cag ctc gag atg acg cta cgc aac tac tac gcg aag cac tgg tgc	432
Ala Gln Leu Glu Met Thr Leu Arg Asn Tyr Tyr Ala Lys His Trp Cys	
130 135 140	
gat ccc tcc ggc ttc gac atc ggt cag cgc gcc ctg tac cag cac ccc	480
Asp Pro Ser Gly Phe Asp Ile Gly Gln Arg Gly Leu Tyr Gln His Pro	
145 150 155 160	
gcg ggg ctg gtg tcc atc ggc gag ttc cag cac ttc aat act ggt gac	528
Ala Gly Leu Val Ser Ile Gly Glu Phe Gln His Phe Asn Thr Gly Asp	
165 170 175	
ccg ctt gac gtc atc atc gat ctc aac atc gtg gcc gag acg gcg ttc	576
Pro Leu Asp Val Ile Ile Asp Leu Asn Ile Val Ala Glu Thr Ala Phe	
180 185 190	
acg aac atc ctg ctg gtg gcc act cca cag gtc gcc gtg gcc aac ggg	624
Thr Asn Ile Leu Leu Val Ala Thr Pro Gln Val Ala Val Ala Asn Gly	
195 200 205	
gac aac gcg atg gcc agc gtg ttc ctc tcg atc cag tcg gac gag gcc	672

Asp Asn Ala Met Ala Ser Val Phe Leu Ser Ile Gln Ser Asp Glu Ala	
210 215 220	
agg cac atg gcc aac ggg tac ggc tcg gtc atg gcg ctg ctg gag aac	720
Arg His Met Ala Asn Gly Tyr Gly Ser Val Met Ala Leu Leu Glu Asn	
225 230 235 240	
gag gac aac ctc ccg ctg ctc aac cag tct ctc gat cgg cac ttc tgg	768
Glu Asp Asn Leu Pro Leu Leu Asn Gln Ser Leu Asp Arg His Phe Trp	
245 250 255	
cgt gcc cac aag gcc ttg gac aac gcg gtc gga tgg tgt tcg gag tat	816
Arg Ala His Lys Ala Leu Asp Asn Ala Val Gly Trp Cys Ser Glu Tyr	
260 265 270	
ggc gcc cgc aag ccg cca tgg agc tac aag gcc cag tgg gag gaa tgg	864
Gly Ala Arg Lys Arg Pro Trp Ser Tyr Lys Ala Gln Trp Glu Glu Trp	
275 280 285	
gtc gtc gac gac ttc gtg ggc ggc tac atc gac cga ctc agc gag ttc	912
Val Val Asp Asp Phe Val Gly Gly Tyr Ile Asp Arg Leu Ser Glu Phe	
290 295 300	
ggc gtt cag gct ccg gcc tgc ctt ggc gcg gcc gcc gac gag gtc aag	960
Gly Val Gln Ala Pro Ala Cys Leu Gly Ala Ala Asp Glu Val Lys	
305 310 315 320	
tgg tcg cac cac acg ctc ggt cag gtg ctg tcg gcg gtg tgg ccg ctg	1008
Trp Ser His His Thr Leu Gly Gln Val Leu Ser Ala Val Trp Pro Leu	
325 330 335	
aac ttc tgg cgc tcg gac gcc atg gga ccg gcg gac ttc gag tgg ttc	1056
Asn Phe Trp Arg Ser Asp Ala Met Gly Pro Ala Asp Phe Glu Trp Phe	
340 345 350	
gag aac cac tac ccg ggc tgg agc gcg gcc tac cag ggt tac tgg gag	1104
Glu Asn His Tyr Pro Gly Trp Ser Ala Ala Tyr Gln Gly Tyr Trp Glu	
355 360 365	
ggc tac aag gcg ctc gcc gac cca gca ggc gga cgc atc atg ctc cag	1152
Gly Tyr Lys Ala Leu Ala Asp Pro Ala Gly Gly Arg Ile Met Leu Gln	
370 375 380	
gag ctg ccg ggt ctg ccg ccg atg tgt cag gtg tgc cag gtg ccg tgc	1200
Glu Leu Pro Gly Leu Pro Pro Met Cys Gln Val Cys Gln Val Pro Cys	
385 390 395 400	
gtg atg ccg ccg ctg gat atg aac gcc gcg ccg atc atc gag ttc gag	1248
Val Met Pro Arg Leu Asp Met Asn Ala Ala Arg Ile Ile Glu Phe Glu	
405 410 415	
ggg cag aaa atc gcg ctg tgc agc gaa ccc tgc cag ccg atc ttc acc	1296
Gly Gln Lys Ile Ala Leu Cys Ser Glu Pro Cys Gln Arg Ile Phe Thr	
420 425 430	
aac tgg ccg gag gcg tac cgc cac cgc aag caa tac tgg gcc cgc tac	1344
Asn Trp Pro Glu Ala Tyr Arg His Arg Lys Gln Tyr Trp Ala Arg Tyr	
435 440 445	
cac gga tgg gac ctg gcg gac gtc atc gtt gat ctc ggc tac atc cgc	1392
His Gly Trp Asp Leu Ala Asp Val Ile Val Asp Leu Gly Tyr Ile Arg	
450 455 460	
ccg gac ggc aag acc ctc atc ggc cag ccg ctg ctc gag atg gag ccg	1440
Pro Asp Gly Lys Thr Leu Ile Gly Gln Pro Leu Leu Glu Met Glu Arg	
465 470 475 480	
ctg tgg acc atc gac gac atc ccg gcc ctt cag tac gaa gtc aag gac	1488
Leu Trp Thr Ile Asp Asp Ile Arg Ala Leu Gln Tyr Glu Val Lys Asp	
485 490 495	
ccg ttg cag gag gcg tga	1506
Pro Leu Gln Glu Ala	
500	

<210> 12
 <211> 501
 <212> PRT
 <213> Nocardia corallina

<400> 12

Met Ala Ser Asn Pro Thr Gln Leu His Glu Lys Ser Lys Ser Tyr Asp
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 Trp Asp Phe Thr Ser Val Glu Arg Arg Pro Lys Phe Glu Thr Lys Tyr
 20 25 30
 Lys Met Pro Lys Lys Gly Lys Asp Pro Phe Arg Val Leu Ile Arg Asp
 35 40 45
 Tyr Met Lys Met Glu Ala Glu Lys Asp Asp Arg Thr His Gly Phe Leu
 50 55 60
 Asp Gly Ala Val Arg Thr Arg Glu Ala Thr Arg Ile Glu Pro Arg Phe
 65 70 75 80
 Ala Glu Ala Met Lys Ile Met Val Pro Gln Leu Thr Asn Ala Glu Tyr
 85 90 95
 Gln Ala Val Ala Gly Cys Gly Met Ile Ile Ser Ala Val Glu Asn Gln
 100 105 110
 Glu Leu Arg Gln Gly Tyr Ala Ala Gln Met Leu Asp Glu Val Arg His
 115 120 125
 Ala Gln Leu Glu Met Thr Leu Arg Asn Tyr Tyr Ala Lys His Trp Cys
 130 135 140
 Asp Pro Ser Gly Phe Asp Ile Gly Gln Arg Gly Leu Tyr Gln His Pro
 145 150 155 160
 Ala Gly Leu Val Ser Ile Gly Glu Phe Gln His Phe Asn Thr Gly Asp
 165 170 175
 Pro Leu Asp Val Ile Ile Asp Leu Asn Ile Val Ala Glu Thr Ala Phe
 180 185 190
 Thr Asn Ile Leu Leu Val Ala Thr Pro Gln Val Ala Val Ala Asn Gly
 195 200 205
 Asp Asn Ala Met Ala Ser Val Phe Leu Ser Ile Gln Ser Asp Glu Ala
 210 215 220
 Arg His Met Ala Asn Gly Tyr Gly Ser Val Met Ala Leu Leu Glu Asn
 225 230 235 240
 Glu Asp Asn Leu Pro Leu Leu Asn Gln Ser Leu Asp Arg His Phe Trp
 245 250 255
 Arg Ala His Lys Ala Leu Asp Asn Ala Val Gly Trp Cys Ser Glu Tyr
 260 265 270
 Gly Ala Arg Lys Arg Pro Trp Ser Tyr Lys Ala Gln Trp Glu Glu Trp
 275 280 285
 Val Val Asp Asp Phe Val Gly Gly Tyr Ile Asp Arg Leu Ser Glu Phe
 290 295 300
 Gly Val Gln Ala Pro Ala Cys Leu Gly Ala Ala Ala Asp Glu Val Lys
 305 310 315 320
 Trp Ser His His Thr Leu Gly Gln Val Leu Ser Ala Val Trp Pro Leu
 325 330 335
 Asn Phe Trp Arg Ser Asp Ala Met Gly Pro Ala Asp Phe Glu Trp Phe
 340 345 350
 Glu Asn His Tyr Pro Gly Trp Ser Ala Ala Tyr Gln Gly Tyr Trp Glu
 355 360 365

Gly Tyr Lys Ala Leu Ala Asp Pro Ala Gly Gly Arg Ile Met Leu Gln
 370 375 380
 Glu Leu Pro Gly Leu Pro Pro Met Cys Gln Val Cys Gln Val Pro Cys
 385 390 395 400
 Val Met Pro Arg Leu Asp Met Asn Ala Ala Arg Ile Ile Glu Phe Glu
 405 410 415
 Gly Gln Lys Ile Ala Leu Cys Ser Glu Pro Cys Gln Arg Ile Phe Thr
 420 425 430
 Asn Trp Pro Glu Ala Tyr Arg His Arg Lys Gln Tyr Trp Ala Arg Tyr
 435 440 445
 His Gly Trp Asp Leu Ala Asp Val Ile Val Asp Leu Gly Tyr Ile Arg
 450 455 460
 Pro Asp Gly Lys Thr Leu Ile Gly Gln Pro Leu Leu Glu Met Glu Arg
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 Leu Trp Thr Ile Asp Asp Ile Arg Ala Leu Gln Tyr Glu Val Lys Asp
 485 490 495
 Pro Leu Gln Glu Ala
 500

<210> 13
 <211> 1494
 <212> DNA
 <213> Xanthobacta sp.

<220>
 <221> CDS
 <222> (1)..(1494)

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 gac tgg acg ctc agc tat gtc gac cgc gcg gtc gcc ttt ccc gag gag 96
 Asp Trp Thr Leu Ser Tyr Val Asp Arg Ala Val Ala Phe Pro Glu Glu
 20 25 30
 tgg aaa ggc gaa aag gac att tgc ggc acg gcc tgg gac gat tgg gac 144
 Trp Lys Gly Glu Lys Asp Ile Cys Gly Thr Ala Trp Asp Asp Trp Asp
 35 40 45
 gag ccc ttc cgg gtc tcc ttc cgc gaa tat gtg atg gtc cag cgc gac 192
 Glu Pro Phe Arg Val Ser Phe Arg Glu Tyr Val Met Val Gln Arg Asp
 50 55 60
 aag gaa gcg agc gtc ggc gcc atc cgc gag gcc atg gtc cgc gcc aag 240
 Lys Glu Ala Ser Val Gly Ala Ile Arg Glu Ala Met Val Arg Ala Lys
 65 70 75 80
 gcc tat gag aag ctc gac gac ggc cac aag gcc acc tcg cac ctg cac 288
 Ala Tyr Glu Lys Leu Asp Asp Gly His Lys Ala Thr Ser His Leu His
 85 90 95
 atg ggc acc atc acc atg gtg gag cac atg gcg gtc acc atg cag agc 336
 Met Gly Thr Ile Thr Met Val Glu His Met Ala Val Thr Met Gln Ser
 100 105 110
 cgg ttc gtg cgc ttc gcg ccg tcc gcc cgc tgg cgc agc ctc ggg gcg 384
 Arg Phe Val Arg Phe Ala Pro Ser Ala Arg Trp Arg Ser Leu Gly Ala
 115 120 125
 ttc ggc atg ctg gac gag acc cgc cac acc cag ctc gac ctg cgc ttc 432
 Phe Gly Met Leu Asp Glu Thr Arg His Thr Gln Leu Asp Leu Arg Phe
 130 135 140
 agc cac gat ctg ctc aac gat tcc ccg agc ttc gac tgg agc cag cgg 480

Ser His Asp Leu Leu Asn Asp Ser Pro Ser Phe Asp Trp Ser Gln Arg	
145 150 155 160	
gcg ttc cac acc gac gaa tgg gcg gtt ctc gcc acc cgc aac ctg ttc	528
Ala Phe His Thr Asp Glu Trp Ala Val Leu Ala Thr Arg Asn Leu Phe	
165 170 175	
gac gac atc atg ctc aac gcc gac tgc gtg gag gcg gcg ctc gcc acc	576
Asp Asp Ile Met Leu Asn Ala Asp Cys Val Glu Ala Ala Leu Ala Thr	
180 185 190	
agc ctg acg ctg gag cac ggc ttc acc aac atc cag ttc gtg gcg ctc	624
Ser Leu Thr Leu Glu His Gly Phe Thr Asn Ile Gln Phe Val Ala Leu	
195 200 205	
gcc tcc gac gcc atg gaa gcc ggc gac gtg aac ttc tcc aac ctc ttg	672
Ala Ser Asp Ala Met Glu Ala Gly Asp Val Asn Phe Ser Asn Leu Leu	
210 215 220	
tcg agc atc cag acc gac gag gcg cgg cac gcc cag ttg ggc ttt ccc	720
Ser Ser Ile Gln Thr Asp Glu Ala Arg His Ala Gln Leu Gly Phe Pro	
225 230 235 240	
acc ctc gac gtg atg atg aag cac gac ccc aag cgc gcc cag cag atc	768
Thr Leu Asp Val Met Met Lys His Asp Pro Lys Arg Ala Gln Gln Ile	
245 250 255	
ctg gac gtc gcc ttc tgg cgc tcc tac cgc atc ttc cag gcg gtg acc	816
Leu Asp Val Ala Phe Trp Arg Ser Tyr Arg Ile Phe Gln Ala Val Thr	
260 265 270	
ggc gtc tcc atg gac tac tac acg ccg gtc gcc aag cgg cag atg tcg	864
Gly Val Ser Met Asp Tyr Tyr Thr Pro Val Ala Lys Arg Gln Met Ser	
275 280 285	
ttc aag gag ttc atg ctg gag tgg atc gtc aag cat cat gag cgc atc	912
Phe Lys Glu Phe Met Leu Glu Trp Ile Val Lys His His Glu Arg Ile	
290 295 300	
ctg cgc gac tac ggc ctc cag aag ccc tgg tac tgg gac acg ttc gag	960
Leu Arg Asp Tyr Gly Leu Gln Lys Pro Trp Tyr Trp Asp Thr Phe Glu	
305 310 315 320	
aag acc ctc gat cac ggc cac cac gcg ctg cac atc ggc acc tgg ttc	1008
Lys Thr Leu Asp His Gly His His Ala Leu His Ile Gly Thr Trp Phe	
325 330 335	
tgg cgc ccg acc ctg ttc tgg gat ccc aat ggc ggc gtc tcg cgc gag	1056
Trp Arg Pro Thr Leu Phe Trp Asp Pro Asn Gly Gly Val Ser Arg Glu	
340 345 350	
gag cgc cgc tgg ctg aac cag aag tat ccg aac tgg gaa gag agc tgg	1104
Glu Arg Arg Trp Leu Asn Gln Lys Tyr Pro Asn Trp Glu Glu Ser Trp	
355 360 365	
ggc gtc ctg tgg gac gag atc atc tcc aac atc aat gcg ggc aac att	1152
Gly Val Leu Trp Asp Glu Ile Ile Ser Asn Ile Asn Ala Gly Asn Ile	
370 375 380	
gaa aag acc ttg ccc gag acg ctg ccg atg ctg tgc aac gtc acc aac	1200
Glu Lys Thr Leu Pro Glu Thr Leu Pro Met Leu Cys Asn Val Thr Asn	
385 390 395 400	
ctg ccc atc ggc tcg cac tgg gac cgc ttc cac ctg aag ccc gag cag	1248
Leu Pro Ile Gly Ser His Trp Asp Arg Phe His Leu Lys Pro Glu Gln	
405 410 415	
ctc gtc tac aag ggg cgc ctc tac acc ttc gac agc gac gtc tcc aag	1296
Leu Val Tyr Lys Gly Arg Leu Tyr Thr Phe Asp Ser Asp Val Ser Lys	
420 425 430	
tgg atc ttc gag ctc gat ccg gag cgc tat gcc ggc cac acc aac gtg	1344
Trp Ile Phe Glu Leu Asp Pro Glu Arg Tyr Ala Gly His Thr Asn Val	
435 440 445	
gtc gac cgc ttc atc ggc ggg cag atc cag ccc atg acc atc gag ggc	1392

Val Asp Arg Phe Ile Gly Gly Gln Ile Gln Pro Met Thr Ile Glu Gly
 450 455 460
 gtg ctc aac tgg atg ggc ctg acg ccc gaa gtc atg ggc aag gac gtg 1440
 Val Leu Asn Trp Met Gly Leu Thr Pro Glu Val Met Gly Lys Asp Val
 465 470 475 480
 ttc aac tac cgt tgg gcc ggc gat tac gcc gag aac cgg atc gcc gcc 1488
 Phe Asn Tyr Arg Trp Ala Gly Asp Tyr Ala Glu Asn Arg Ile Ala Ala
 485 490 495
 gag taa 1494
 Glu

<210> 14
 <211> 497
 <212> PRT
 <213> Xanthobacta sp.

<400> 14

Met Ala Leu Leu Asn Arg Asp Asp Trp Tyr Asp Ile Ala Arg Asp Val
 1 5 10 15
 Asp Trp Thr Leu Ser Tyr Val Asp Arg Ala Val Ala Phe Pro Glu Glu
 20 25 30
 Trp Lys Gly Glu Lys Asp Ile Cys Gly Thr Ala Trp Asp Asp Trp Asp
 35 40 45
 Glu Pro Phe Arg Val Ser Phe Arg Glu Tyr Val Met Val Gln Arg Asp
 50 55 60
 Lys Glu Ala Ser Val Gly Ala Ile Arg Glu Ala Met Val Arg Ala Lys
 65 70 75 80
 Ala Tyr Glu Lys Leu Asp Asp Gly His Lys Ala Thr Ser His Leu His
 85 90 95
 Met Gly Thr Ile Thr Met Val Glu His Met Ala Val Thr Met Gln Ser
 100 105 110
 Arg Phe Val Arg Phe Ala Pro Ser Ala Arg Trp Arg Ser Leu Gly Ala
 115 120 125
 Phe Gly Met Leu Asp Glu Thr Arg His Thr Gln Leu Asp Leu Arg Phe
 130 135 140
 Ser His Asp Leu Leu Asn Asp Ser Pro Ser Phe Asp Trp Ser Gln Arg
 145 150 155 160
 Ala Phe His Thr Asp Glu Trp Ala Val Leu Ala Thr Arg Asn Leu Phe
 165 170 175
 Asp Asp Ile Met Leu Asn Ala Asp Cys Val Glu Ala Ala Leu Ala Thr
 180 185 190
 Ser Leu Thr Leu Glu His Gly Phe Thr Asn Ile Gln Phe Val Ala Leu
 195 200 205
 Ala Ser Asp Ala Met Glu Ala Gly Asp Val Asn Phe Ser Asn Leu Leu 210
 215 220
 Ser Ser Ile Gln Thr Asp Glu Ala Arg His Ala Gln Leu Gly Phe Pro
 225 230 235 240
 Thr Leu Asp Val Met Met Lys His Asp Pro Lys Arg Ala Gln Gln Ile
 245 250 255
 Leu Asp Val Ala Phe Trp Arg Ser Tyr Arg Ile Phe Gln Ala Val Thr
 260 265 270
 Gly Val Ser Met Asp Tyr Tyr Thr Pro Val Ala Lys Arg Gln Met Ser
 275 280 285

Phe Lys Glu Phe Met Leu Glu Trp Ile Val Lys His His Glu Arg Ile
 290 295 300
 Leu Arg Asp Tyr Gly Leu Gln Lys Pro Trp Tyr Trp Asp Thr Phe Glu
 305 310 315 320
 Lys Thr Leu Asp His Gly His His Ala Leu His Ile Gly Thr Trp Phe
 325 330 335
 Trp Arg Pro Thr Leu Phe Trp Asp Pro Asn Gly Gly Val Ser Arg Glu
 340 345 350
 Glu Arg Arg Trp Leu Asn Gln Lys Tyr Pro Asn Trp Glu Glu Ser Trp
 355 360 365
 Gly Val Leu Trp Asp Glu Ile Ile Ser Asn Ile Asn Ala Gly Asn Ile
 370 375 380
 Glu Lys Thr Leu Pro Glu Thr Leu Pro Met Leu Cys Asn Val Thr Asn
 385 390 395 400
 Leu Pro Ile Gly Ser His Trp Asp Arg Phe His Leu Lys Pro Glu Gln
 405 410 415
 Leu Val Tyr Lys Gly Arg Leu Tyr Thr Phe Asp Ser Asp Val Ser Lys
 420 425 430
 Trp Ile Phe Glu Leu Asp Pro Glu Arg Tyr Ala Gly His Thr Asn Val
 435 440 445
 Val Asp Arg Phe Ile Gly Gly Gln Ile Gln Pro Met Thr Ile Glu Gly
 450 455 460
 Val Leu Asn Trp Met Gly Leu Thr Pro Glu Val Met Gly Lys Asp Val
 465 470 475 480
 Phe Asn Tyr Arg Trp Ala Gly Asp Tyr Ala Glu Asn Arg Ile Ala Ala
 485 490 495
 Glu

<210> 15
 <211> 1026
 <212> DNA
 <213> Xanthobacta sp.

<220>
 <221> CDS
 <222> (1)..(1026)

<400> 15
 atg aca cag cag cgc ccc acc cgc acg cgc gag cgc aag aag acc tgg 48
 Met Thr Gln Gln Arg Pro Thr Arg Thr Arg Glu Arg Lys Lys Thr Trp
 1 5 10 15
 acg gct ttc ggc aat ctc gga cgc aag ccg acc gac tac gag gtc gtc 96
 Thr Ala Phe Gly Asn Leu Gly Arg Lys Pro Thr Asp Tyr Glu Val Val
 20 25 30
 acc cac aac atg aac cac acc atg cgc gcc acg ccc ctg gag ctg tcg 144
 Thr His Asn Met Asn His Thr Met Arg Gly Thr Pro Leu Glu Leu Ser
 35 40 45
 ccg acg gtg cac gcc aat gtg tgg ctc aag aag aac cgc gac gag atc 192
 Pro Thr Val His Ala Asn Val Trp Leu Lys Lys Asn Arg Asp Glu Ile
 50 55 60
 gcg ctc aag gtc gac agc tgg gat ctg ttc cgc gat ccc gac cgc acc 240
 Ala Leu Lys Val Asp Ser Trp Asp Leu Phe Arg Asp Pro Asp Arg Thr
 65 70 75 80
 acc tac gac acc tac gtc aag atg cag gac gac cag gag acc tat gtc 288

Thr Tyr Asp Thr Tyr Val Lys Met Gln Asp Asp Gln Glu Thr Tyr Val
 85 90 95
 gac aac ctg ctc ctg tcc tac acc ggc gag ggc cgc tac gac gag gag 336
 Asp Asn Leu Leu Leu Ser Tyr Thr Gly Glu Gly Arg Tyr Asp Glu Glu
 100 105 110
 ctt tcc tcg cgc agc ctc gac ctc ctg tcc gcg ggg ctg acg ccg acc 384
 Leu Ser Ser Arg Ser Leu Asp Leu Leu Ser Ala Gly Leu Thr Pro Thr
 115 120 125
 cgc tat ctg ggc cat ggg ctg cag atg ctc gcg gcc tat atc cag cag 432
 Arg Tyr Leu Gly His Gly Leu Gln Met Leu Ala Ala Tyr Ile Gln Gln
 130 135 140
 ctc gcc ccg tcg gcc tat gtg ggc aat tgc gcg gtg ttc cag acc tcc 480
 Leu Ala Pro Ser Ala Tyr Val Gly Asn Cys Ala Val Phe Gln Thr Ser
 145 150 155 160
 gac gcg ctg cgc cgc gtg cag cgc gtc gcc tac cgc acc cgc cag ctc 528
 Asp Ala Leu Arg Arg Val Gln Arg Val Ala Tyr Arg Thr Arg Gln Leu
 165 170 175
 gcc gac gcc cat ccg gcc cgc ggc ttc ggc tcc ggc gac cgg gcg gtg 576
 Ala Asp Ala His Pro Ala Arg Gly Phe Gly Ser Gly Asp Arg Ala Val
 180 185 190
 tgg gag aag tcc ccg gac tgg cag ccc atc cgc aag gcc atc gag gag 624
 Trp Glu Lys Ser Pro Asp Trp Gln Pro Ile Arg Lys Ala Ile Glu Glu
 195 200 205
 ctg ctc gtc acc ttc gaa tgg gac aag gcg ctc gcc ggc acc aat ttc 672
 Leu Leu Val Thr Phe Glu Trp Asp Lys Ala Leu Ala Gly Thr Asn Phe
 210 215 220
 gtg gtg aag ccg atc ctc gac gag ctg ttc ctc aac cac ctg gcg cgc 720
 Val Val Lys Pro Ile Leu Asp Glu Leu Phe Leu Asn His Leu Ala Arg
 225 230 235 240
 ctg ctc cac gtg gag ggc gac gag ctc gac agc ctc gtg ctg cgg aac 768
 Leu Leu His Val Glu Gly Asp Glu Leu Asp Ser Leu Val Leu Arg Asn
 245 250 255
 ctt cac ggc gac gcc cag cgc cac gcc cgc tgg acg gcc gcg ctc ggc 816
 Leu His Gly Asp Ala Gln Arg His Ala Arg Trp Thr Ala Ala Leu Gly
 260 265 270
 cgc ttc gcc gtc gag cag aac gtg aac aac cgc acg gtc ctg cgc gac 864
 Arg Phe Ala Val Glu Gln Asn Val Asn Asn Arg Thr Val Leu Arg Asp
 275 280 285
 gcc atc gcc ggc tgg cac gag acc ggc gag gcg gtc ctc gcc gcg ggc 912
 Ala Ile Ala Gly Trp His Glu Thr Gly Glu Ala Val Leu Ala Ala Gly
 290 295 300
 gcc ggg atg ctt gcg agc cgc gcc ccc agc gcg gat gcg gcc aag atc 960
 Ala Gly Met Leu Ala Ser Arg Ala Pro Ser Ala Asp Ala Ala Lys Ile
 305 310 315 320
 gcc gac gag gtc cgc gcc acg ctc gcg cag ctg cac gcc aat gcg ggc 1008
 Ala Asp Glu Val Arg Ala Thr Leu Ala Gln Leu His Ala Asn Ala Gly
 325 330 335
 ctc ggg cac gat gcc tga 1026
 Leu Gly His Asp Ala
 340
 <210> 16
 <211> 341
 <212> PRT
 <213> Xanthobacta sp.
 <400> 16
 Met Thr Gln Gln Arg Pro Thr Arg Thr Arg Glu Arg Lys Lys Thr Trp

1	5	10	15
Thr Ala Phe Gly Asn Leu Gly Arg Lys Pro Thr Asp Tyr Glu Val Val	20	25	30
Thr His Asn Met Asn His Thr Met Arg Gly Thr Pro Leu Glu Leu Ser	35	40	45
Pro Thr Val His Ala Asn Val Trp Leu Lys Lys Asn Arg Asp Glu Ile	50	55	60
Ala Leu Lys Val Asp Ser Trp Asp Leu Phe Arg Asp Pro Asp Arg Thr	65	70	75
Thr Tyr Asp Thr Tyr Val Lys Met Gln Asp Asp Gln Glu Thr Tyr Val	85	90	95
Asp Asn Leu Leu Leu Ser Tyr Thr Gly Glu Gly Arg Tyr Asp Glu Glu	100	105	110
Leu Ser Ser Arg Ser Leu Asp Leu Leu Ser Ala Gly Leu Thr Pro Thr	115	120	125
Arg Tyr Leu Gly His Gly Leu Gln Met Leu Ala Ala Tyr Ile Gln Gln	130	135	140
Leu Ala Pro Ser Ala Tyr Val Gly Asn Cys Ala Val Phe Gln Thr Ser	145	150	155
Asp Ala Leu Arg Arg Val Gln Arg Val Ala Tyr Arg Thr Arg Gln Leu	165	170	175
Ala Asp Ala His Pro Ala Arg Gly Phe Gly Ser Gly Asp Arg Ala Val	180	185	190
Trp Glu Lys Ser Pro Asp Trp Gln Pro Ile Arg Lys Ala Ile Glu Glu	195	200	205
Leu Leu Val Thr Phe Glu Trp Asp Lys Ala Leu Ala Gly Thr Asn Phe	210	215	220
Val Val Lys Pro Ile Leu Asp Glu Leu Phe Leu Asn His Leu Ala Arg	225	230	235
Leu Leu His Val Glu Gly Asp Glu Leu Asp Ser Leu Val Leu Arg Asn	245	250	255
Leu His Gly Asp Ala Gln Arg His Ala Arg Trp Thr Ala Ala Leu Gly	260	265	270
Arg Phe Ala Val Glu Gln Asn Val Asn Asn Arg Thr Val Leu Arg Asp	275	280	285
Ala Ile Ala Gly Trp His Glu Thr Gly Glu Ala Val Leu Ala Ala Gly	290	295	300
Ala Gly Met Leu Ala Ser Arg Ala Pro Ser Ala Asp Ala Ala Lys Ile	305	310	315
Ala Asp Glu Val Arg Ala Thr Leu Ala Gln Leu His Ala Asn Ala Gly	325	330	335
Leu Gly His Asp Ala	340		

<210> 17
 <211> 267
 <212> DNA
 <213> Xanthobacta sp.

<220>
 <221> CDS
 <222> (1)..(267)

<400> 17
 atg tct ttg ttc ccc atc gtg ggc cgc ttc gtg ggg gat ttc gtc ccc 48
 Met Ser Leu Phe Pro Ile Val Gly Arg Phe Val Gly Asp Phe Val Pro
 1 5 10 15
 cac ctg gtg gcg gtg gac acc tct gac acc atc gat cag atc gcc gag 96
 His Leu Val Ala Val Asp Thr Ser Asp Thr Ile Asp Gln Ile Ala Glu
 20 25 30
 aag gtg gcg gtc cac acg gtc ggg cgg cgc ttg ccg ccc gat ccc acc 144
 Lys Val Ala Val His Thr Val Gly Arg Arg Leu Pro Pro Asp Pro Thr
 35 40 45
 gcc acc ggc tat gag gtg ctc ctc gac ggc gag acc ctg gac ggg ggc 192
 Ala Thr Gly Tyr Glu Val Ile Met Thr Lys Arg Glu Thr Leu Asp Gly Gly
 50 55 60
 gcc acc ctg gag gcc atc atg acc aag cgc gag atg ctg ccc ctg cag 240
 Ala Thr Leu Glu Ala Ile Met Thr Lys Arg Glu Met Leu Pro Leu Gln
 65 70 75 80
 tgg ttc gac gtg agg ttc aag aag tga 267
 Trp Phe Asp Val Arg Phe Lys Lys
 85

<210> 18
 <211> 88
 <212> PRT
 <213> Xanthobacta sp.

<400> 18
 Met Ser Leu Phe Pro Ile Val Gly Arg Phe Val Gly Asp Phe Val Pro
 1 5 10 15
 His Leu Val Ala Val Asp Thr Ser Asp Thr Ile Asp Gln Ile Ala Glu
 20 25 30
 Lys Val Ala Val His Thr Val Gly Arg Arg Leu Pro Pro Asp Pro Thr
 35 40 45
 Ala Thr Gly Tyr Glu Val Leu Leu Asp Gly Glu Thr Leu Asp Gly Gly
 50 55 60
 Ala Thr Leu Glu Ala Ile Met Thr Lys Arg Glu Met Leu Pro Leu Gln
 65 70 75 80
 Trp Phe Asp Val Arg Phe Lys Lys
 85

<210> 19
 <211> 1584
 <212> DNA
 <213> Methylococcus capsulatas

<220>
 <221> CDS
 <222> (1)..(1584)

<400> 19
 atg gca ctt agc acc gca acc aag gcc gcg acg gac gcg ctg gct gcc 48
 Met Ala Leu Ser Thr Ala Thr Lys Ala Ala Thr Asp Ala Leu Ala Ala
 1 5 10 15
 aat cgg gca ccc acc agc gtg aat gca cag gaa gtg cac cgt tgg ctc 96
 Asn Arg Ala Pro Thr Ser Val Asn Ala Gln Glu Val His Arg Trp Leu
 20 25 30
 cag agc ttc aac tgg gat ttc aag aac aac cgg acc aag tac gcc acc 144
 Gln Ser Phe Asn Trp Asp Phe Lys Asn Asn Arg Thr Lys Tyr Ala Thr
 35 40 45
 aag tac aag atg gcg aac gag acc aag gaa cag ttc aag ctg atc gcc 192

Lys 50	Tyr	Lys	Met	Ala	Asn	Glu	Thr	Lys	Glu	Gln	Phe	Lys	Leu	Ile	Ala	
aag	gaa	tat	gcg	cgc	atg	gag	gca	gtc	aag	gac	gaa	agg	cag	ttc	ggt	240
Lys 65	Glu	Tyr	Ala	Arg	Met	Glu	Ala	Val	Lys	Asp	Glu	Arg	Gln	Phe	Gly	80
agc	ctg	cag	gat	gcg	ctg	acc	cgc	ctc	aac	gcc	ggt	gtt	cgc	gtt	cat	288
Ser	Leu	Gln	Asp	Ala	Leu	Thr	Arg	Leu	Asn	Ala	Gly	Val	Arg	Val	His	95
ccg	aag	tgg	aac	gag	acc	atg	aaa	gtg	gtt	tcg	aac	ttc	ctg	gaa	gtg	336
Pro	Lys	Trp	Asn	Glu	Thr	Met	Lys	Val	Val	Ser	Asn	Phe	Leu	Glu	Val	110
ggc	gaa	tac	aac	gcc	atc	gcc	gct	acc	ggg	atg	ctg	tgg	gat	tcc	gcc	384
Gly	Glu	Tyr	Asn	Ala	Ile	Ala	Ala	Thr	Gly	Met	Leu	Trp	Asp	Ser	Ala	125
cag	gcg	gcg	gaa	cag	aag	aac	ggc	tat	ctg	gcc	cag	gtg	ttg	gat	gaa	432
Gln	Ala	Ala	Glu	Gln	Lys	Asn	Gly	Tyr	Leu	Ala	Gln	Val	Leu	Asp	Glu	140
atc	cgc	cac	acc	cac	cag	tgt	gcc	tac	gtc	aac	tac	tac	ttc	gcg	aag	480
Ile	Arg	His	Thr	His	Gln	Cys	Ala	Tyr	Val	Asn	Tyr	Tyr	Phe	Ala	Lys	155
aac	ggc	cag	gac	ccg	gcc	ggt	cac	aac	gat	gct	cgc	cgc	acc	cgt	acc	528
Asn	Gly	Gln	Asp	Pro	Ala	Gly	His	Asn	Asp	Ala	Arg	Arg	Thr	Arg	Thr	175
atc	ggt	ccg	ctg	tgg	aag	ggc	atg	aag	cgc	gtg	ttt	tcc	gac	ggc	ttc	576
Ile	Gly	Pro	Leu	Trp	Lys	Gly	Met	Lys	Arg	Val	Phe	Ser	Asp	Gly	Phe	185
att	tcc	ggc	gac	gcc	gtg	gaa	tgc	tcc	ctc	aac	ctg	cag	ctg	gtg	ggt	624
Ile	Ser	Gly	Asp	Ala	Val	Glu	Cys	Ser	Leu	Asn	Leu	Gln	Leu	Val	Gly	195
gag	gcc	tgc	ttc	acc	aat	ccg	ctg	atc	gtc	gca	gtg	acc	gaa	tgg	gct	672
Glu	Ala	Cys	Phe	Thr	Asn	Pro	Leu	Ile	Val	Ala	Val	Thr	Glu	Trp	Ala	210
gcc	gcc	aac	ggc	gat	gaa	atc	acc	ccg	acg	gtg	ttc	ctg	tcg	atc	gag	720
Ala	Ala	Asn	Gly	Asp	Glu	Ile	Thr	Pro	Thr	Val	Phe	Leu	Ser	Ile	Glu	225
acc	gac	gaa	ctg	cgc	cac	atg	gcc	aac	ggt	tac	cag	acc	gtc	gtt	tcc	768
Thr	Asp	Glu	Leu	Arg	His	Met	Ala	Asn	Gly	Tyr	Gln	Thr	Val	Val	Ser	245
atc	gcc	aac	gat	ccg	gct	tcc	gcc	aag	tat	ctc	aac	acg	gac	ctg	aac	816
Ile	Ala	Asn	Asp	Pro	Ala	Ser	Ala	Lys	Tyr	Leu	Asn	Thr	Asp	Leu	Asn	260
aac	gcc	ttc	tgg	acc	cag	cag	aag	tac	ttc	acg	ccg	gtg	ttg	ggc	atg	864
Asn	Ala	Phe	Trp	Thr	Gln	Gln	Lys	Tyr	Phe	Thr	Pro	Val	Leu	Gly	Met	275
ctg	ttc	gag	tat	ggc	tcc	aag	ttc	aag	gtc	gag	ccg	tgg	gtc	aag	acg	912
Leu	Phe	Glu	Tyr	Gly	Ser	Lys	Phe	Lys	Val	Glu	Pro	Trp	Val	Lys	Thr	290
tgg	gac	cgc	tgg	gtg	tac	gag	gac	tgg	ggc	ggc	atc	tgg	atc	ggc	cgt	960
Trp	Asp	Arg	Trp	Val	Tyr	Glu	Asp	Trp	Gly	Gly	Ile	Trp	Ile	Gly	Arg	305
ctg	ggc	aag	tac	ggg	gtg	gag	tcg	ccg	cgc	agc	ctc	aag	gac	gcc	aag	1008
Leu	Gly	Lys	Tyr	Gly	Val	Glu	Ser	Pro	Arg	Ser	Leu	Lys	Asp	Ala	Lys	325
cag	gac	gct	tac	tgg	gct	cac	cac	gac	ctg	tat	ctg	ctg	gct	tat	gcg	1056
Gln	Asp	Ala	Tyr	Trp	Ala	His	His	Asp	Leu	Tyr	Leu	Leu	Ala	Tyr	Ala	340
ctg	tgg	ccg	acc	ggc	ttc	ttc	cgt	ctg	gcg	ctg	ccg	gat	cag	gaa	gaa	1104

Leu Trp Pro Thr Gly Phe Phe Arg Leu Ala Leu Pro Asp Gln Glu Glu
 355 360 365
 atg gag tgg ttc gag gcc aac tac ccc ggc tgg tac gac cac tac ggc 1152
 Met Glu Trp Phe Glu Ala Asn Tyr Pro Gly Trp Tyr Asp His Tyr Gly
 370 375 380
 aag atc tac gag gaa tgg cgc gcc cgc ggt tgc gag gat ccg tcc tcg 1200
 Lys Ile Tyr Glu Glu Trp Arg Ala Arg Gly Cys Glu Asp Pro Ser Ser
 385 390 395 400
 ggc ttc atc ccg ctg atg tgg ttc atc gaa aac aac cat ccc atc tac 1248
 Gly Phe Ile Pro Leu Met Trp Phe Ile Glu Asn Asn His Pro Ile Tyr
 405 410 415
 atc gat cgc gtg tgc caa gtg ccg ttc tgc ccg agc ttg gcc aag ggc 1296
 Ile Asp Arg Val Ser Gln Val Pro Phe Cys Pro Ser Leu Ala Lys Gly
 420 425 430
 gcc agc acc ctg cgc gtg cac gag tac aac ggc gag atg cac acc ttc 1344
 Ala Ser Thr Leu Arg Val His Glu Tyr Asn Gly Glu Met His Thr Phe
 435 440 445
 agc gac cag tgg ggc gag cgc atg tgg ctg gcc gag ccg gag cgc tac 1392
 Ser Asp Gln Trp Gly Glu Arg Met Trp Leu Ala Glu Pro Glu Arg Tyr
 450 455 460
 gag tgc cag aac atc ttc gaa cag tac gaa gga cgc gaa ctg tcg gaa 1440
 Glu Cys Gln Asn Ile Phe Glu Gln Tyr Glu Gly Arg Glu Leu Ser Glu
 465 470 475 480
 gtg atc gcc gaa ctg cac ggg ctg cgc agt gat ggc aag acc ctg atc 1488
 Val Ile Ala Glu Leu His Gly Leu Arg Ser Asp Gly Lys Thr Leu Ile
 485 490 495
 gcc cag ccg cat gtc cgt ggc gac aag ctg tgg acg ttg gac gat atc 1536
 Ala Gln Pro His Val Arg Gly Asp Lys Leu Trp Thr Leu Asp Asp Ile
 500 505 510
 aaa cgc ctg aac tgc gtc ttc aag aac ccg gtg aag gca ttc aat tga 1584
 Lys Arg Leu Asn Cys Val Phe Lys Asn Pro Val Lys Ala Phe Asn
 515 520 525

<210> 20
 <211> 527
 <212> PRT
 <213> Methylococcus capsulatas

<400> 20

Met Ala Leu Ser Thr Ala Thr Lys Ala Ala Thr Asp Ala Leu Ala Ala
 1 5 10 15
 Asn Arg Ala Pro Thr Ser Val Asn Ala Gln Glu Val His Arg Trp Leu
 20 25 30
 Gln Ser Phe Asn Trp Asp Phe Lys Asn Asn Arg Thr Lys Tyr Ala Thr
 35 40 45
 Lys Tyr Lys Met Ala Asn Glu Thr Lys Glu Gln Phe Lys Leu Ile Ala
 50 55 60
 Lys Glu Tyr Ala Arg Met Glu Ala Val Lys Asp Glu Arg Gln Phe Gly
 65 70 75 80
 Ser Leu Gln Asp Ala Leu Thr Arg Leu Asn Ala Gly Val Arg Val His
 85 90 95
 Pro Lys Trp Asn Glu Thr Met Lys Val Val Ser Asn Phe Leu Glu Val
 100 105 110
 Gly Glu Tyr Asn Ala Ile Ala Ala Thr Gly Met Leu Trp Asp Ser Ala
 115 120 125
 Gln Ala Ala Glu Gln Lys Asn Gly Tyr Leu Ala Gln Val Leu Asp Glu

130	135	140
Ile Arg His Thr His Gln Cys Ala Tyr Val Asn Tyr Tyr Phe Ala Lys 145 150 155 160		
Asn Gly Gln Asp Pro Ala Gly His Asn Asp Ala Arg Arg Thr Arg Thr 165 170 175		
Ile Gly Pro Leu Trp Lys Gly Met Lys Arg Val Phe Ser Asp Gly Phe 180 185 190		
Ile Ser Gly Asp Ala Val Glu Cys Ser Leu Asn Leu Gln Leu Val Gly 195 200 205		
Glu Ala Cys Phe Thr Asn Pro Leu Ile Val Ala Val Thr Glu Trp Ala 210 215 220		
Ala Ala Asn Gly Asp Glu Ile Thr Pro Thr Val Phe Leu Ser Ile Glu 225 230 235 240		
Thr Asp Glu Leu Arg His Met Ala Asn Gly Tyr Gln Thr Val Val Ser 245 250 255		
Ile Ala Asn Asp Pro Ala Ser Ala Lys Tyr Leu Asn Thr Asp Leu Asn 260 265 270		
Asn Ala Phe Trp Thr Gln Gln Lys Tyr Phe Thr Pro Val Leu Gly Met 275 280 285		
Leu Phe Glu Tyr Gly Ser Lys Phe Lys Val Glu Pro Trp Val Lys Thr 290 295 300		
Trp Asp Arg Trp Val Tyr Glu Asp Trp Gly Gly Ile Trp Ile Gly Arg 305 310 315 320		
Leu Gly Lys Tyr Gly Val Glu Ser Pro Arg Ser Leu Lys Asp Ala Lys 325 330 335		
Gln Asp Ala Tyr Trp Ala His His Asp Leu Tyr Leu Leu Ala Tyr Ala 340 345 350		
Leu Trp Pro Thr Gly Phe Phe Arg Leu Ala Leu Pro Asp Gln Glu Glu 355 360 365		
Met Glu Trp Phe Glu Ala Asn Tyr Pro Gly Trp Tyr Asp His Tyr Gly 370 375 380		
Lys Ile Tyr Glu Glu Trp Arg Ala Arg Gly Cys Glu Asp Pro Ser Ser 385 390 395 400		
Gly Phe Ile Pro Leu Met Trp Phe Ile Glu Asn Asn His Pro Ile Tyr 405 410 415		
Ile Asp Arg Val Ser Gln Val Pro Phe Cys Pro Ser Leu Ala Lys Gly 420 425 430		
Ala Ser Thr Leu Arg Val His Glu Tyr Asn Gly Glu Met His Thr Phe 435 440 445		
Ser Asp Gln Trp Gly Glu Arg Met Trp Leu Ala Glu Pro Glu Arg Tyr 450 455 460		
Glu Cys Gln Asn Ile Phe Glu Gln Tyr Glu Gly Arg Glu Leu Ser Glu 465 470 475 480		
Val Ile Ala Glu Leu His Gly Leu Arg Ser Asp Gly Lys Thr Leu Ile 485 490 495		
Ala Gln Pro His Val Arg Gly Asp Lys Leu Trp Thr Leu Asp Asp Ile 500 505 510		
Lys Arg Leu Asn Cys Val Phe Lys Asn Pro Val Lys Ala Phe Asn 515 520 525		

<211> 1170
 <212> DNA
 <213> Methylococcus capsulatas

<220>
 <221> CDS
 <222> (1)..(1170)

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<400> 21
atg agc atg tta gga gaa aga cgc cgc ggt ctg acc gat ccg gaa atg      48
Met Ser Met Leu Gly Glu Arg Arg Arg Gly Leu Thr Asp Pro Glu Met
1          5          10          15

gcg gcc gtc att ttg aag gcg ctt cct gaa gct ccg ctg gac ggc aac      96
Ala Ala Val Ile Leu Lys Ala Leu Pro Glu Ala Pro Leu Asp Gly Asn
20          25          30

aac aag atg ggt tat ttc gtc acc ccc cgc tgg aaa cgc ttg acg gaa      144
Asn Lys Met Gly Tyr Phe Val Thr Pro Arg Trp Lys Arg Leu Thr Glu
35          40          45

tat gaa gcc ctg acc gtt tat gcg cag ccc aac gcc gac tgg atc gcc      192
Tyr Glu Ala Leu Thr Val Tyr Ala Gln Pro Asn Ala Asp Trp Ile Ala
50          55          60

ggc ggc ctg gac tgg ggc gac tgg acc cag aaa ttc cac ggc ggc cgc      240
Gly Gly Leu Asp Trp Gly Asp Trp Thr Gln Lys Phe His Gly Gly Arg
65          70          75

cct tcc tgg ggc aac gag acc acg gag ctg cgc acc gtc gac tgg ttc      288
Pro Ser Trp Gly Asn Glu Thr Thr Glu Leu Arg Thr Val Asp Trp Phe
85          90          95

aag cac cgt gac ccg ctc cgc cgt tgg cat gcg ccg tac gtc aag gac      336
Lys His Arg Asp Pro Leu Arg Arg Trp His Ala Pro Tyr Val Lys Asp
100         105         110

aag gcc gag gaa tgg cgc tac acc gac cgc ttc ctg cag ggt tac tcc      384
Lys Ala Glu Trp Arg Tyr Thr Asp Arg Phe Leu Gln Gly Tyr Ser
115         120         125

gcc gac ggt cag atc cgg gcg atg aac ccg acc tgg cgg gac gag ttc      432
Ala Asp Gly Gln Ile Arg Ala Met Asn Pro Thr Trp Arg Asp Glu Phe
130         135         140

atc aac cgg tat tgg ggc gcc ttc ctg ttc aac gaa tac gga ttg ttc      480
Ile Asn Arg Tyr Trp Gly Ala Phe Leu Phe Asn Glu Tyr Gly Leu Phe
145         150         155

aac gct cat tcg cag ggc gcc cgg gag gcg ctg tcg gac gta acc cgc      528
Asn Ala His Ser Gln Gly Ala Arg Glu Ala Leu Ser Asp Val Thr Arg
165         170         175

gtc agc ctg gct ttc tgg ggc ttc gac aag atc gac atc gcc cag atg      576
Val Ser Leu Ala Phe Trp Gly Phe Asp Lys Ile Asp Ile Ala Gln Met
180         185         190

atc caa ctc gaa cgg ggt ttc ctc gcc aag atc gta ccc ggt ttc gac      624
Ile Gln Leu Glu Arg Gly Phe Leu Ala Lys Ile Val Pro Gly Phe Asp
195         200         205

gag tcc aca gcg gtg ccg aag gcc gaa tgg acg aac ggg gag gtc tac      672
Glu Ser Thr Ala Val Pro Lys Ala Glu Trp Thr Asn Gly Glu Val Tyr
210         215         220

aag agc gcc cgt ctg gcc gtg gaa ggg ctg tgg cag gag gtg ttc gac      720
Lys Ser Ala Arg Leu Ala Val Glu Gly Leu Trp Gln Glu Val Phe Asp
225         230         235         240

tgg aac gag agc gct ttc tcg gtg cac gcc gtc tat gac gcg ctg ttc      768
Trp Asn Glu Ser Ala Phe Ser Val His Ala Val Tyr Asp Ala Leu Phe
245         250         255

ggt cag ttc gtc cgc cgc gag ttc ttt cag cgg ctg gct ccc cgc ttc      816
Gly Gln Phe Val Arg Arg Glu Phe Phe Gln Arg Leu Ala Pro Arg Phe

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260	265	270	
ggc gac aat ctg acg cca ttc ttc atc aac cag gcc cag aca tac ttc Gly Asp Asn Leu Thr Pro Phe Phe Ile Asn Gln Ala Gln Thr Tyr Phe 275 280 285			864
cag atc gcc aag cag ggc gta cag gat ctg tat tac aac tgt ctg ggt Gln Ile Ala Lys Gln Gly Val Gln Asp Leu Tyr Tyr Asn Cys Leu Gly 290 295 300			912
gac gat ccg gag ttc agc gat tac aac cgt acc gtg atg cgc aac tgg Asp Asp Pro Glu Phe Ser Asp Tyr Asn Arg Thr Val Met Arg Asn Trp 305 310 315 320			960
acc ggc aag tgg ctg gag ccc acg atc gcc gct ctg cgc gac ttc atg Thr Gly Lys Trp Leu Glu Pro Thr Ile Ala Ala Leu Arg Asp Phe Met 325 330 335			1008
ggg ctg ttt gcg aag ctg ccg gcg ggc acc act gac aag gaa gaa atc Gly Leu Phe Ala Lys Leu Pro Ala Gly Thr Thr Asp Lys Glu Glu Ile 340 345 350			1056
acc gcg tcc ctg tac ccg gtg gtc gac gac tgg atc gag gac tac gcc Thr Ala Ser Leu Tyr Arg Val Val Asp Asp Trp Ile Glu Asp Tyr Ala 355 360 365			1104
agc gcg atc gac ttc aag gcg gac cgc gat cag atc gtt aaa gcg gtt Ser Ala Ile Asp Phe Lys Ala Asp Arg Asp Gln Ile Val Lys Ala Val 370 375 380			1152
ctg gca gga ttg aaa taa Leu Ala Gly Leu Lys 385			1170
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Met Ser Met Leu Gly Glu Arg Arg Arg Gly Leu Thr Asp Pro Glu Met 1 5 10 15			
Ala Ala Val Ile Leu Lys Ala Leu Pro Glu Ala Pro Leu Asp Gly Asn 20 25 30			
Asn Lys Met Gly Tyr Phe Val Thr Pro Arg Trp Lys Arg Leu Thr Glu 35 40 45			
Tyr Glu Ala Leu Thr Val Tyr Ala Gln Pro Asn Ala Asp Trp Ile Ala 50 55 60			
Gly Gly Leu Asp Trp Gly Asp Trp Thr Gln Lys Phe His Gly Gly Arg 65 70 75 80			
Pro Ser Trp Gly Asn Glu Thr Thr Glu Leu Arg Thr Val Asp Trp Phe 85 90 95			
Lys His Arg Asp Pro Leu Arg Arg Trp His Ala Pro Tyr Val Lys Asp 100 105 110			
Lys Ala Glu Glu Trp Arg Tyr Thr Asp Arg Phe Leu Gln Gly Tyr Ser 115 120 125			
Ala Asp Gly Gln Ile Arg Ala Met Asn Pro Thr Trp Arg Asp Glu Phe 130 135 140			
Ile Asn Arg Tyr Trp Gly Ala Phe Leu Phe Asn Glu Tyr Gly Leu Phe 145 150 155 160			
Asn Ala His Ser Gln Gly Ala Arg Glu Ala Leu Ser Asp Val Thr Arg 165 170 175			
Val Ser Leu Ala Phe Trp Gly Phe Asp Lys Ile Asp Ile Ala Gln Met			

180 185 190

Ile Gln Leu Glu Arg Gly Phe Leu Ala Lys Ile Val Pro Gly Phe Asp
195 200 205

Glu Ser Thr Ala Val Pro Lys Ala Glu Trp Thr Asn Gly Glu Val Tyr
210 215 220

Lys Ser Ala Arg Leu Ala Val Glu Gly Leu Trp Gln Glu Val Phe Asp
225 230 235 240

Trp Asn Glu Ser Ala Phe Ser Val His Ala Val Tyr Asp Ala Leu Phe
245 250 255

Gly Gln Phe Val Arg Arg Glu Phe Phe Gln Arg Leu Ala Pro Arg Phe
260 265 270

Gly Asp Asn Leu Thr Pro Phe Phe Ile Asn Gln Ala Gln Thr Tyr Phe
275 280 285

Gln Ile Ala Lys Gln Gly Val Gln Asp Leu Tyr Tyr Asn Cys Leu Gly
290 295 300

Asp Asp Pro Glu Phe Ser Asp Tyr Asn Arg Thr Val Met Arg Asn Trp
305 310 315 320

Thr Gly Lys Trp Leu Glu Pro Thr Ile Ala Ala Leu Arg Asp Phe Met
325 330 335

Gly Leu Phe Ala Lys Leu Pro Ala Gly Thr Thr Asp Lys Glu Glu Ile
340 345 350

Thr Ala Ser Leu Tyr Arg Val Val Asp Asp Trp Ile Glu Asp Tyr Ala
355 360 365

Ser Ala Ile Asp Phe Lys Ala Asp Arg Asp Gln Ile Val Lys Ala Val
370 375 380

Leu Ala Gly Leu Lys
385

<210> 23
<211> 513
<212> DNA
<213> Methylococcus capsulatas

<220>
<221> CDS
<222> (1)..(513)

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Met Ala Lys Leu Gly Ile His Ser Asn Asp Thr Arg Asp Ala Trp Val
1 5 10 15

96
aac aag atc gcg cag ctc aac acc ctg gaa aaa gcg gcc gag atg ctg
Asn Lys Ile Ala Gln Leu Asn Thr Leu Glu Lys Ala Ala Glu Met Leu
20 25 30

144
aag cag ttc cgg atg gac cac acc acg ccg ttc cgc aac agc tac gaa
Lys Gln Phe Arg Met Asp His Thr Thr Pro Phe Arg Asn Ser Tyr Glu
35 40 45

192
ctg gac aac gac tac ctc tgg atc gag gcc aag ctc gaa gag aag gtc
Leu Asp Asn Asp Tyr Leu Trp Ile Glu Ala Lys Leu Glu Glu Lys Val
50 55 60

240
gcc gtc ctc aag gca cgc gcc ttc aac gag gtg gac ttc cgt cat aag
Ala Val Leu Lys Ala Arg Ala Phe Asn Glu Val Asp Phe Arg His Lys
65 70 75 80

288
acc gct ttc ggc gag gat gcc aag tcc gtt ctg gac ggc acc gtc gcg
Thr Ala Phe Gly Glu Asp Ala Lys Ser Val Leu Asp Gly Thr Val Ala
85 90 95

aag atg aac gcg gcc aag gac aag tgg gag gcg gag aag atc cat atc 336
 Lys Met Asn Ala Ala Lys Asp Lys Trp Glu Ala Glu Lys Ile His Ile
 100 105 110

ggt ttc cgc cag gcc tac aag ccg ccg atc atg ccg gtg aac tat ttc 384
 Gly Phe Arg Gln Ala Tyr Lys Pro Pro Ile Met Pro Val Asn Tyr Phe
 115 120 125

ctg gac ggc gag cgt cag ttg ggg acc ccg ctg atg gaa ctg cgc aac 432
 Leu Asp Gly Glu Arg Gln Leu Gly Thr Arg Leu Met Glu Leu Arg Asn
 130 135 140

ctc aac tac tac gac acg ccg ctg gaa gaa ctg cgc aaa cag cgc ggt 480
 Leu Asn Tyr Tyr Asp Thr Pro Leu Glu Glu Leu Arg Lys Gln Arg Gly
 145 150 155 160

gtg cgg gtg gtg cat ctg cag tgg ccg cac tga 513
 Val Arg Val Val His Leu Gln Ser Pro His
 165 170

<210> 24
 <211> 170
 <212> PRT
 <213> *Methylococcus capsulatas*

<400> 24

Met Ala Lys Leu Gly Ile His Ser Asn Asp Thr Arg Asp Ala Trp Val
 1 5 10 15

Asn Lys Ile Ala Gln Leu Asn Thr Leu Glu Lys Ala Ala Glu Met Leu
 20 25 30

Lys Gln Phe Arg Met Asp His Thr Thr Pro Phe Arg Asn Ser Tyr Glu
 35 40 45

Leu Asp Asn Asp Tyr Leu Trp Ile Glu Ala Lys Leu Glu Glu Lys Val
 50 55 60

Ala Val Leu Lys Ala Arg Ala Phe Asn Glu Val Asp Phe Arg His Lys
 65 70 75 80

Thr Ala Phe Gly Glu Asp Ala Lys Ser Val Leu Asp Gly Thr Val Ala
 85 90 95

Lys Met Asn Ala Ala Lys Asp Lys Trp Glu Ala Glu Lys Ile His Ile
 100 105 110

Gly Phe Arg Gln Ala Tyr Lys Pro Pro Ile Met Pro Val Asn Tyr Phe
 115 120 125

Leu Asp Gly Glu Arg Gln Leu Gly Thr Arg Leu Met Glu Leu Arg Asn
 130 135 140

Leu Asn Tyr Tyr Asp Thr Pro Leu Glu Glu Leu Arg Lys Gln Arg Gly
 145 150 155 160

Val Arg Val Val His Leu Gln Ser Pro His
 165 170

<210> 25
 <211> 1206
 <212> DNA
 <213> *Pseudomonas oleovorans*

<220>
 <221> CDS
 <222> (1)..(1206)

<400> 25

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 Met Leu Glu Lys His Arg Val Leu Asp Ser Ala Pro Glu Tyr Val Asp

1	5	10	15	
aaa aag aaa tat ctc tgg ata cta tca act ttg tgg ccg gct act ccg				96
Lys Lys Lys Tyr Leu Trp Ile Leu Ser Thr Leu Trp Pro Ala Thr Pro	20	25	30	
atg atc gga atc tgg ctt gca aat gaa act ggt tgg ggg att ttt tat				144
Met Ile Gly Ile Trp Leu Ala Asn Glu Thr Gly Trp Gly Ile Phe Tyr	35	40	45	
ggg ctg gta ttg ctc gta tgg tac ggc gca ctt cca ttg ctt gat gcg				192
Gly Leu Val Leu Leu Val Trp Tyr Gly Ala Leu Pro Leu Leu Asp Ala	50	55	60	
atg ttt ggt gag gac ttt aat aat ccg cct gaa gaa gtg gtg ccg aaa				240
Met Phe Gly Glu Asp Phe Asn Asn Pro Pro Glu Glu Val Val Pro Lys	65	70	75	80
cta gag aag gag cgg tac tat cga gtt ttg aca tat cta aca gtt cct				288
Leu Glu Lys Glu Arg Tyr Tyr Arg Val Leu Thr Tyr Leu Thr Val Pro	85	90	95	
atg cat tac gct gca tta att gtg tca gca tgg tgg gtc gga act cag				336
Met His Tyr Ala Ala Leu Ile Val Ser Ala Trp Trp Val Gly Thr Gln	100	105	110	
cca atg tct tgg ctt gaa att ggt gcg ctt gcc ttg tca ctg ggt atc				384
Pro Met Ser Trp Leu Glu Ile Gly Ala Leu Ala Leu Ser Leu Gly Ile	115	120	125	
gtg aac gga cta gcg ctc aat aca gga cac gaa ctc ggt cac aag aag				432
Val Asn Gly Leu Ala Leu Asn Thr Gly His Glu Leu Gly His Lys Lys	130	135	140	
gag act ttt gat cgt tgg atg gcc aaa att gtg ttg gct gtc gta ggg				480
Glu Thr Phe Asp Arg Trp Met Ala Lys Ile Val Leu Ala Val Val Gly	145	150	155	160
tac ggt cac ttc ttt att gag cat aat aag ggt cat cac cgt gat gtc				528
Tyr Gly His Phe Phe Ile Glu His Asn Lys Gly His His Arg Asp Val	165	170	175	
gct aca ccg atg gat cct gca aca tcc ccg atg gga gaa agc att tat				576
Ala Thr Pro Met Asp Pro Ala Thr Ser Arg Met Gly Glu Ser Ile Tyr	180	185	190	
aag ttt tca atc cgt gag atc cca gga gca ttt att cgt gct tgg ggg				624
Lys Phe Ser Ile Arg Glu Ile Pro Gly Ala Phe Ile Arg Ala Trp Gly	195	200	205	
ctt gag gaa caa cgc ctt tcg cgc cgt ggc caa agc gtt tgg agt ttc				672
Leu Glu Glu Gln Arg Leu Ser Arg Arg Gly Gln Ser Val Trp Ser Phe	210	215	220	
gat aat gaa atc ctc caa cca atg atc atc aca gtt att ctt tac gcc				720
Asp Asn Glu Ile Leu Gln Pro Met Ile Ile Thr Val Ile Leu Tyr Ala	225	230	235	240
gtt ctc ctt gcc ttg ttt gga cct aag atg ctg gtg ttc ctg ccg att				768
Val Leu Leu Ala Leu Phe Gly Pro Lys Met Leu Val Phe Leu Pro Ile	245	250	255	
caa atg gct ttc ggt tgg tgg cag ctg acc agt gcg aac tat att gaa				816
Gln Met Ala Phe Gly Trp Trp Gln Leu Thr Ser Ala Asn Tyr Ile Glu	260	265	270	
cat tac ggc ttg ctc cgt caa aaa atg gag gac ggt cga tat gag cat				864
His Tyr Gly Leu Leu Arg Gln Lys Met Glu Asp Gly Arg Tyr Glu His	275	280	285	
caa aag ccg cac cat tct tgg aat agt aat cac atc gtc tct aat cta				912
Gln Lys Pro His His Ser Trp Asn Ser Asn His Ile Val Ser Asn Leu	290	295	300	
gtg ctg ttc cac ctt cag ccg cac tcg gat cac cac gcg cat cca aca				960
Val Leu Phe His Leu Gln Arg His Ser Asp His His Ala His Pro Thr				

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<210> 26
<211> 401
<212> PRT
<213> Pseudomonas oleovorans
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Asp Asn Glu Ile Leu Gln Pro Met Ile Ile Thr Val Ile Leu Tyr Ala
 225 230 235 240

Val Leu Leu Ala Leu Phe Gly Pro Lys Met Leu Val Phe Leu Pro Ile
 245 250 255

Gln Met Ala Phe Gly Trp Trp Gln Leu Thr Ser Ala Asn Tyr Ile Glu
 260 265 270

His Tyr Gly Leu Leu Arg Gln Lys Met Glu Asp Gly Arg Tyr Glu His
 275 280 285

Gln Lys Pro His His Ser Trp Asn Ser Asn His Ile Val Ser Asn Leu
 290 295 300

Val Leu Phe His Leu Gln Arg His Ser Asp His His Ala His Pro Thr
 305 310 315 320

Arg Ser Tyr Gln Ser Leu Arg Asp Phe Pro Gly Leu Pro Ala Leu Pro
 325 330 335

Thr Gly Tyr Pro Gly Ala Phe Leu Met Ala Met Ile Pro Gln Trp Phe
 340 345 350

Arg Ser Val Met Asp Pro Lys Val Val Asp Trp Ala Gly Gly Asp Leu
 355 360 365

Asn Lys Ile Gln Ile Asp Asp Ser Met Arg Glu Thr Tyr Leu Lys Lys
 370 375 380

Phe Gly Thr Ser Ser Ala Gly His Ser Ser Ser Thr Ser Ala Val Ala
 385 390 395 400

Ser

<210> 27
 <211> 1560
 <212> DNA
 <213> Burkholderia cepacia

<220>
 <221> CDS
 <222> (1)..(1560)

<400> 27
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 Met Asp Thr Ser Val Gln Lys Lys Lys Leu Gly Leu Lys Asn Arg Tyr
 1 5 10 15

gca gcg atg acc cgc ggt ctt ggc tgg cag acc agc tac cag ccg atg 96
 Ala Ala Met Thr Arg Gly Leu Gly Trp Gln Thr Ser Tyr Gln Pro Met
 20 25 30

gag aaa gtg ttt ccg tac gac aag tac gaa ggc atc aag atc cac gat 144
 Glu Lys Val Phe Pro Tyr Asp Lys Tyr Glu Gly Ile Lys Ile His Asp
 35 40 45

tgg gat aaa tgg gaa gac ccc ttc cgc ctg acc atg gac gcc tac tgg 192
 Trp Asp Lys Trp Glu Asp Pro Phe Arg Leu Thr Met Asp Ala Tyr Trp
 50 55 60

aaa tat cag ggc gag aag gaa aaa aag ctt tac gcc gtc atc gac gct 240
 Lys Tyr Gln Gly Glu Lys Glu Lys Lys Leu Tyr Ala Val Ile Asp Ala
 65 70 75 80

ttc gcg cag aac aac ggg cag ttg agc att tcc gac gcg cga tat gtc 288
 Phe Ala Gln Asn Asn Gly Gln Leu Ser Ile Ser Asp Ala Arg Tyr Val
 85 90 95

aac gca ctc aag gtg ttt atc cag ggt gtg aca ccg ttg gag tat atg 336
 Asn Ala Leu Lys Val Phe Ile Gln Gly Val Thr Pro Leu Glu Tyr Met
 100 105 110

gca cac cga ggt ttt gcc cac att ggt cgg cat ttt acg ggt gaa ggg Ala His Arg Gly Phe Ala His Ile Gly Arg His Phe Thr Gly Glu Gly 115 120 125	384
gca cgt gtt gct tgc cag atg cag tcc atc gac gag ctg cgt cac ttc Ala Arg Val Ala Cys Gln Met Gln Ser Ile Asp Glu Leu Arg His Phe 130 135 140	432
cag acc gaa atg cat gct ctc tcg cac tac aac aag tat ttt aac ggt Gln Thr Glu Met His Ala Leu Ser His Tyr Asn Lys Tyr Phe Asn Gly 145 150 155 160	480
ctg cac aac tcc atc cat tgg tac gac cgg gtt tgg tat ttg tcg gtg Leu His Asn Ser Ile His Trp Tyr Asp Arg Val Trp Tyr Leu Ser Val 165 170 175	528
ccc aag tca ttt ttt gaa gac gcg gcc acc ggt gga ccg ttc gag ttt Pro Lys Ser Phe Phe Glu Asp Ala Ala Thr Gly Gly Pro Phe Glu Phe 180 185 190	576
ctt acc gcg gtg agc ttt tcg ttc gaa tat gtg ttg acc aac ctg ctg Leu Thr Ala Val Ser Phe Ser Phe Glu Tyr Val Leu Thr Asn Leu Leu 195 200 205	624
ttt gtc ccc ttc atg tcg ggt gct gct tac aac ggg gac atg tct acg Phe Val Pro Phe Met Ser Gly Ala Ala Tyr Asn Gly Asp Met Ser Thr 210 215 220	672
gtc act ttc ggt ttt tcg gcg caa agt gac gaa tcg cgc cac atg aca Val Thr Phe Gly Phe Ser Ala Gln Ser Asp Glu Ser Arg His Met Thr 225 230 235 240	720
ctc gcc atc gaa tgc atc aag ttc atg cta gaa cag gat ccg gac aac Leu Gly Ile Glu Cys Ile Lys Phe Met Leu Glu Gln Asp Pro Asp Asn 245 250 255	768
gtg ccc atc gtg cag cgc tgg atc gac aag tgg ttc tgg cgc ggc tat Val Pro Ile Val Gln Arg Trp Ile Asp Lys Trp Phe Trp Arg Gly Tyr 260 265 270	816
cgg ctg ttg agc atc gtg gcc atg atg cag gac tac atg ctg ccc aac Arg Leu Leu Ser Ile Val Ala Met Met Gln Asp Tyr Met Leu Pro Asn 275 280 285	864
cgg gtg atg agc tgg cgc gag agc tgg gag atg tac gtc gag cag aac Arg Val Met Ser Trp Arg Glu Ser Trp Glu Met Tyr Val Glu Gln Asn 290 295 300	912
ggc ggc gcg ctg ttc aag gat ctt gcg cgt tat ggc atc cgc aag ccc Gly Gly Ala Leu Phe Lys Asp Leu Ala Arg Tyr Gly Ile Arg Lys Pro 305 310 315 320	960
aag ggc tgg gac cag gct tgc gaa ggc aag gac cac atc agc cat cag Lys Gly Trp Asp Gln Ala Cys Glu Gly Lys Asp His Ile Ser His Gln 325 330 335	1008
acc ttc gcc gta ttc tat aac tat aac gcc gcg gcc ccc atc cac acc Thr Phe Ala Val Phe Tyr Asn Tyr Asn Ala Ala Ala Pro Ile His Thr 340 345 350	1056
tgg gtt ccc aca aaa gaa gaa atg gga tgg ctg tcg gag aag tac ccc Trp Val Pro Thr Lys Glu Glu Met Gly Trp Leu Ser Glu Lys Tyr Pro 355 360 365	1104
gag acg ttc gac aag tat tac cgt ccg cgt tgg gac tac tgg cgc gag Glu Thr Phe Asp Lys Tyr Tyr Arg Pro Arg Trp Asp Tyr Trp Arg Glu 370 375 380	1152
cag gcc gcc aag ggc aac cgt ttc tac aac aag acg ctg ccg atg ctc Gln Ala Ala Lys Gly Asn Arg Phe Tyr Asn Lys Thr Leu Pro Met Leu 385 390 395 400	1200
tgc act acc tgc cag att ccg atg ata ttc acc gag cct ggc gac gca Cys Thr Thr Cys Gln Ile Pro Met Ile Phe Thr Glu Pro Gly Asp Ala 405 410 415	1248

acc aag atc tgc tat cgc gag tgc gcc tac ctc ggc gac aag tat cac 1296
 Thr Lys Ile Cys Tyr Arg Glu Ser Ala Tyr Leu Gly Asp Lys Tyr His
 420 425 430

ttc tgc agc gac cac tgc aag gag att ttt gac aac gaa ccc gaa aag 1344
 Phe Cys Ser Asp His Cys Lys Glu Ile Phe Asp Asn Glu Pro Glu Lys
 435 440 445

ttc gtg cag tca tgg ctt ccg ccg cag caa gtg tat caa gga aac tgt 1392
 Phe Val Gln Ser Trp Leu Pro Pro Gln Gln Val Tyr Gln Gly Asn Cys
 450 455 460

ttc aag ccg gat gcc gat ccg acc aag gag ggt ttt gat ccc ttg atg 1440
 Phe Lys Pro Asp Ala Asp Pro Thr Lys Glu Gly Phe Asp Pro Leu Met
 465 470 475 480

gcc ttg ctc gac tac tac aac ctg aat gta ggc cgg gac aac ttc gat 1488
 Ala Leu Leu Asp Tyr Tyr Asn Leu Asn Val Gly Arg Asp Asn Phe Asp
 485 490 495

ttc gag gga tgc gaa gac caa aag aac ttt gct gcc tgg cgt gga gag 1536
 Phe Glu Gly Ser Glu Asp Gln Lys Asn Phe Ala Ala Trp Arg Gly Glu
 500 505 510

gtc ttg caa gga gaa gcc aaa tga 1560
 Val Leu Gln Gly Glu Ala Lys
 515

<210> 28
 <211> 519
 <212> PRT
 <213> Burkholderia cepacia

<400> 28

Met Asp Thr Ser Val Gln Lys Lys Lys Leu Gly Leu Lys Asn Arg Tyr
 1 5 10 15

Ala Ala Met Thr Arg Gly Leu Gly Trp Gln Thr Ser Tyr Gln Pro Met
 20 25 30

Glu Lys Val Phe Pro Tyr Asp Lys Tyr Glu Gly Ile Lys Ile His Asp
 35 40 45

Trp Asp Lys Trp Glu Asp Pro Phe Arg Leu Thr Met Asp Ala Tyr Trp
 50 55 60

Lys Tyr Gln Gly Glu Lys Glu Lys Lys Leu Tyr Ala Val Ile Asp Ala
 65 70 75 80

Phe Ala Gln Asn Asn Gly Gln Leu Ser Ile Ser Asp Ala Arg Tyr Val
 85 90 95

Asn Ala Leu Lys Val Phe Ile Gln Gly Val Thr Pro Leu Glu Tyr Met
 100 105 110

Ala His Arg Gly Phe Ala His Ile Gly Arg His Phe Thr Gly Glu Gly
 115 120 125

Ala Arg Val Ala Cys Gln Met Gln Ser Ile Asp Glu Leu Arg His Phe
 130 135 140

Gln Thr Glu Met His Ala Leu Ser His Tyr Asn Lys Tyr Phe Asn Gly
 145 150 155 160

Leu His Asn Ser Ile His Trp Tyr Asp Arg Val Trp Tyr Leu Ser Val
 165 170 175

Pro Lys Ser Phe Phe Glu Asp Ala Ala Thr Gly Gly Pro Phe Glu Phe
 180 185 190

Leu Thr Ala Val Ser Phe Ser Phe Glu Tyr Val Leu Thr Asn Leu Leu
 195 200 205

Phe Val Pro Phe Met Ser Gly Ala Ala Tyr Asn Gly Asp Met Ser Thr
 210 215 220
 Val Thr Phe Gly Phe Ser Ala Gln Ser Asp Glu Ser Arg His Met Thr
 225 230 235 240
 Leu Gly Ile Glu Cys Ile Lys Phe Met Leu Glu Gln Asp Pro Asp Asn
 245 250 255
 Val Pro Ile Val Gln Arg Trp Ile Asp Lys Trp Phe Trp Arg Gly Tyr
 260 265 270
 Arg Leu Leu Ser Ile Val Ala Met Met Gln Asp Tyr Met Leu Pro Asn
 275 280 285
 Arg Val Met Ser Trp Arg Glu Ser Trp Glu Met Tyr Val Glu Gln Asn
 290 295 300
 Gly Gly Ala Leu Phe Lys Asp Leu Ala Arg Tyr Gly Ile Arg Lys Pro
 305 310 315 320
 Lys Gly Trp Asp Gln Ala Cys Glu Gly Lys Asp His Ile Ser His Gln
 325 330 335
 Thr Phe Ala Val Phe Tyr Asn Tyr Asn Ala Ala Ala Pro Ile His Thr
 340 345 350
 Trp Val Pro Thr Lys Glu Glu Met Gly Trp Leu Ser Glu Lys Tyr Pro
 355 360 365
 Glu Thr Phe Asp Lys Tyr Tyr Arg Pro Arg Trp Asp Tyr Trp Arg Glu
 370 375 380
 Gln Ala Ala Lys Gly Asn Arg Phe Tyr Asn Lys Thr Leu Pro Met Leu
 385 390 395 400
 Cys Thr Thr Cys Gln Ile Pro Met Ile Phe Thr Glu Pro Gly Asp Ala
 405 410 415
 Thr Lys Ile Cys Tyr Arg Glu Ser Ala Tyr Leu Gly Asp Lys Tyr His
 420 425 430
 Phe Cys Ser Asp His Cys Lys Glu Ile Phe Asp Asn Glu Pro Glu Lys
 435 440 445
 Phe Val Gln Ser Trp Leu Pro Pro Gln Gln Val Tyr Gln Gly Asn Cys
 450 455 460
 Phe Lys Pro Asp Ala Asp Pro Thr Lys Glu Gly Phe Asp Pro Leu Met
 465 470 475 480
 Ala Leu Leu Asp Tyr Tyr Asn Leu Asn Val Gly Arg Asp Asn Phe Asp
 485 490 495
 Phe Glu Gly Ser Glu Asp Gln Lys Asn Phe Ala Ala Trp Arg Gly Glu
 500 505 510
 Val Leu Gln Gly Glu Ala Lys
 515

<210> 29
 <211> 996
 <212> DNA
 <213> Burkholderia cepacia

<220>
 <221> CDS
 <222> (1)..(996)

<400> 29
 atg acc atc gat ttg aag acg cgg gaa atc aaa cca ctg cgt cac acc
 Met Thr Ile Asp Leu Lys Thr Arg Glu Ile Lys Pro Leu Arg His Thr
 1 5 10 15

tac acg cac gtg gct caa tac atc ggg gcc gat aaa gcc gct tcg cgc Tyr Thr His Val Ala Gln Tyr Ile Gly Ala Asp Lys Ala Ala Ser Arg 20 25 30	96
tat cag gaa ggc act gta ggt gct caa ccc gca gcg aat ttt cat tac Tyr Gln Glu Gly Thr Val Gly Ala Gln Pro Ala Ala Asn Phe His Tyr 35 40 45	144
cgg ccc acg tgg gat ccc gag cat gaa ctg ttc gac acg tcg cgt acc Arg Pro Thr Trp Asp Pro Glu His Glu Leu Phe Asp Thr Ser Arg Thr 50 55 60	192
gcg att caa atg aag gac tgg tat gcg ctg aaa gac ccg cgt cag ttc Ala Ile Gln Met Lys Asp Trp Tyr Ala Leu Lys Asp Pro Arg Gln Phe 65 70 75 80	240
tac tac gcg tcg tgg acg atg acc cga gcg cgg cag caa gac gcg atg Tyr Tyr Ala Ser Trp Thr Met Thr Arg Ala Arg Gln Gln Asp Ala Met 85 90 95	288
gaa tcc aac ttc gag ttt gtc gag tcg cgc ggc atg atc gat ctc gtt Glu Ser Asn Phe Glu Phe Val Glu Ser Arg Gly Met Ile Asp Leu Val 100 105 110	336
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cac gcg gcc tgg ggc gcg aac atg aac aac tcc cag atc tgt gcc cta His Ala Ala Trp Gly Ala Asn Met Asn Asn Ser Gln Ile Cys Ala Leu 130 135 140	432
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gga ccc gat ctt ctt gac gaa gcc aag caa gcc tgg atg acg agt ccc Gly Pro Asp Leu Leu Asp Glu Ala Lys Gln Ala Trp Met Thr Ser Pro 180 185 190	576
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 Arg Pro Thr Trp Asp Pro Glu His Glu Leu Phe Asp Thr Ser Arg Thr
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 Ala Ile Gln Met Lys Asp Trp Tyr Ala Leu Lys Asp Pro Arg Gln Phe
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 Tyr Tyr Ala Ser Trp Thr Met Thr Arg Ala Arg Gln Gln Asp Ala Met
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 Gly Pro Asp Leu Leu Asp Glu Ala Lys Gln Ala Trp Met Thr Ser Pro
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 195 200 205
 Asp Pro Val Glu Leu Phe Ile Ala Gln Asn Leu Ala Leu Asp Gly Leu
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 225 230 235 240
 Asn Gly Gly Ser Ala Val Ala Met Leu Thr Thr Phe Met Pro Glu Trp
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 His Asp Glu Ser Ser Arg Trp Val Asp Ala Val Val Lys Thr Met Ala
 260 265 270
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 275 280 285
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 Gly Gln Pro Trp His Pro Asp Pro Ala Lys Ser Leu Ala Glu Asn Gly
 85 90 95
 ctg acg cac aaa gac gtg atc cgc ttt cgc acg cct ggc ttg aac ggt 336
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 Phe Gly Ala Leu Cys Thr Gln Val Leu Pro Gly Thr Tyr Gly Tyr His
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 Pro Asp Phe Ser Lys Ile Asp Trp Ser Gln Val Gln Trp Phe Lys Ser
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 Gly Gln Pro Trp His Pro Asp Pro Ala Lys Ser Leu Ala Glu Asn Gly
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Leu Leu Ala Lys Ala Glu Glu Ile Gly Arg Ile Ala Glu Glu Glu Ala
20      25      30

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195      200      205

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cag ctt gag gct tgc caa aaa gaa gga aag acg gtg atg aac gat atg 912
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gag cga gag cag cta ttc gca tgg cgt gga tat gtg gca aaa gcg tcc 960
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Gln Leu Glu Ala Cys Gln Lys Glu Gly Lys Thr Val Met Asn Asp Met 290 295 300			
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Ala Asn Ile Ala Val Arg Thr Leu Leu Thr Leu Gly Gly Asn Ser Ile 325 330 335			
Phe Lys Gly Asp Pro Val Glu Leu Phe Thr Arg Asp Leu Leu Ala Val 340 345 350			
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aag ccc ttt acc cct cca agg gag gtt cac caa cag gtg cta cac tca Lys Pro Phe Thr Pro Pro Arg Glu Val His Gln Gln Val Leu His Ser 50 55 60			192
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gaa aat aac ata ttg gtt cac cta aag cct gtc gaa aaa tgc tgg caa Glu Asn Asn Ile Leu Val His Leu Lys Pro Val Glu Lys Cys Trp Gln 85 90 95			288
gca cag gat ttc cta cca gat ccc gca tct gac gga ttt atg gaa caa Ala Gln Asp Phe Leu Pro Asp Pro Ala Ser Asp Gly Phe Met Glu Gln 100 105 110			336
gtg gag gaa tta cgg gct cgg gct aag gag att ccg gat gat tac ttt			384

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 145 150 155 160
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 Thr Leu Leu Leu Gly Leu Val Trp Thr Arg Ala Trp Thr Ala Glu Glu
 165 170 175
 aac agg cac ggt gat ctt cta cat cag tat ctg tat ctt agt ggg cgg 576
 Asn Arg His Gly Asp Leu Leu His Gln Tyr Leu Tyr Leu Ser Gly Arg
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 Val Asp Met Arg Gln Ile Gln Lys Thr Ile Gln Tyr Leu Ile Gly Ser
 195 200 205
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 Gly Met Asp Pro Arg Thr Glu Asn Ser Pro Tyr Leu Gly Phe Ile Tyr
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 Ile Ile Ala Ala Asp Glu Lys Arg His Glu Thr Ala Tyr Thr Lys Ile
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 370 375 380
 Pro Phe Ser Trp Ile Phe Asp Arg Glu Val Lys Leu
 385 390 395

CLAIMS

1. A method of carrying out an oxidation reaction catalysed by a monooxygenase enzyme and using hydrogen peroxide as an oxidant, in which reaction a low level of oxidation damage of the monooxygenase occurs, said method comprising producing the hydrogen peroxide simultaneously with the oxidation reaction, wherein the hydrogen peroxide is produced at a rate less than or equal to the rate at which it is used in the reaction.

2. A method according to claim 1, wherein the monooxygenase enzyme has a K_m for H_2O_2 of at least 15nM.

3. A method according to claim 1 or 2, wherein the monooxygenase enzyme is a P450 enzyme.

4. A method according to any one of the preceding claims, wherein the rate of H_2O_2 production is less than or equal to 3 μg per mg of enzyme.

5. A method according to any one of the preceding claims, wherein the concentration of H_2O_2 throughout the reaction is less than or equal to 1 mM.

6. A method according to any one of the preceding claims, wherein the reaction continues for at least 240 minutes.

7. A method according to any one of the preceding claims, wherein the H_2O_2 is produced by an electrochemical reaction.

8. A method according to any one of claims 1 to 6, wherein the H_2O_2 is produced by an enzyme reaction.

9. A method according to claim 8, wherein the enzyme is glucose oxidase.

10. A method according to any one of claims 1 to 6, wherein the H_2O_2 is
5 produced by a H_2O_2 precursor.

11. A method according to claim 10, wherein the H_2O_2 precursor is perborate, percarbonate or perphosphate.

12. A method according to any one of the preceding claims, wherein
10 the substrate which is oxidised by the monooxygenase enzyme is an alkane, aromatic compound, terpenoid compound, alkene or fatty acid.

13. Use of electrodes for producing H_2O_2 to drive an oxidation reaction
15 as defined in claim 7.

14. Use of an enzyme for producing H_2O_2 to drive an oxidation reaction as defined in claim 8 or 9.

15. Use of perborate, percarbonate or perphosphate for producing H_2O_2
20 to drive an oxidation reaction as defined in claim 10.

16. A method of carrying out an oxidation reaction catalysed by a monooxygenase enzyme and using hydrogen peroxide as an oxidant, in which
25 reaction a low level of oxidation damage of the monooxygenase occurs, said method comprising carrying out the reaction in the presence of an H_2O_2 or hydroxyl radical sequestering agent that controls the H_2O_2 or hydroxyl radical concentration.

17. A method according to claim 16, wherein the sequestering agent is
30 EDTA.

